97055022 Rev. 00 2019-03

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1. FOREWORD

The instructions inform the user on how to properly operate the device. It is extremely important to read this manual carefully and thoroughly before using the device.

This publication must not be reproduced, copied or transferred in any manner (electronically, mechanically, via photocopies, translations or other means) without the prior written consent of the manufacturer.

The manufacturer has a company policy of continual development. Therefore, some of the instructions, specifications and figures given in this manual may slightly differ from the purchased product. The manufacturer reserves the right to make changes to this manual without giving prior notice. The original text is in Italian; this is a translation from the original in Italian.

1.1. SYMBOLS USED



NOTE:

Pay particular attention to the paragraphs marked with the symbol shown.



CAUTION:

Potential danger for people, environment and property. Follow the procedures indicated in the manual to prevent potential damage to materials, devices and/or property.

1.2. SYMBOLS ON THE DEVICE



Potential danger due to high temperature.



Disposal symbol in accordance with Directive 2012/19/EU.



Device in compliance with essential requirements of Directive 93/42/EU and subsequent modifications.

Notified body: IMQ spa



Refer to the user manual.



Device compliant with the requirements set out in the Directive 2014/68/EU (PED), category II, for 22 and 28 I sterilizers.

Notified body: CSI spa



Ukrainian national symbol of conformity.



Device compliant with the requirements set out in the Directive 2014/68/EU (PED), category I, for 17 I sterilizers.

Notified body: CSI spa

1.3. RELEVANT EUROPEAN DIRECTIVES

The product described in this manual is manufactured in accordance with the highest safety standards and doesn't represent any danger for the operator if used according to the following instructions. The product is **complying** with the following **European Directives as applicable**:

93/42/EEC, and subsequent amendments and additions, concerning medical devices.

2011/65/EU, (Rohs II) on restriction of hazardous substances in electrical and electronic devices.

The product complies with Standard EN 13060:2014.

1.4. CLASSIFICATION

Classification of the device according to the rules indicated in Annex IX of Directive 93/42/EEC and subsequent modifications and integrations: **CLASS IIB**.

1.5. INTENDED USE

The product described in this manual is only intended for sterilization of reusable surgical instruments and materials.

DEVICE INTENDED FOR PROFESSIONAL USE



The use of the device is strictly reserved to qualified personnel. It must never be used or handled by untrained and/or unauthorised persons.

The device must not be used for the sterilization of fluids, liquids or pharmaceutical products.



The sterilizer is not a mobile or portable device

1.5.1. IMPORTANT NOTES



Information contained in this manual are subject to change without notice.

The manufacturer is not responsible for direct, indirect or accidental damage resulting from or relating to the provision or use of this information. This document may not be reproduced, adapted or translated, in part or in full, without the prior written permission of the manufacturer.

1.6. GENERAL WARNINGS

When using this product, always follow the instructions in the manual and never use it for anything other than its intended purpose.



The user is responsible for any legal requirements relating to the installation and use of the product. The manufacturer will not be held responsible for any breakage, malfunction, property damage or injury to people in the event that the product is not installed or used correctly, or proper maintenance is not carried out.

Please observe the following precautions in order to avoid injury or property damage:

· Use ONLY demineralised and/or distilled water of high quality (IF THE DEMINERALISATION FILTER IS NOT PRESENT IN THE FILLING TANK).



The use of water of inadequate quality can severely damage the device. See technical characteristics appendix in this regard.

- · Do not pour water or other fluids on the device;
- · Do not pour flammable substances on the device;
- · Do not use the system in the presence of flammable or explosive gases or vapours;
- · Before performing any maintenance or cleaning intervention, ALWAYS DISCONNECT power supply.



Whenever it is not possible to disconnect the power supply from the device, or if the external mains switch is distant or not visible to the maintenance technician, place a "work in progress" sign on the external mains switch after having turned it off.



- Make sure the electrical system is grounded according to current laws and/or standards;
- Do not remove any label or nameplate from the device; request new ones, if necessary.
- Use only original spare parts.







1.7. RESIDUAL RISKS

FOR THE USER

- · Contamination due to improper handling of the load;
- · Burn by contact with hot surfaces or fluids.

FOR THE PATIENT

- · Contamination due to unsterilised material caused by wrong cleaning treatment before sterilization;
- · Contamination due to implementation of a wrong reuse process;
- Contamination due to material unsuitable to sterilization or not compliant with instructions for use;
- · Contamination due to unsterilised material caused by wrong final assessment of sterilization process;
- · Contamination due to missing or wrong scheduled maintenance;
- · Contamination due to missing periodic validation.

1.8. INFORMATION ON MITIGATION OF RESIDUAL RISKS

FOR THE USER

Contamination due to improper handling of the load.

See chapter PREPARING THE MATERIAL.

Burn by contact with hot surfaces or fluids.

To extract the sterile material, once the sterilization process has been completed with saturated steam at 121° or 134°, proceed as follows:

- · Always wear PPE suitable for the handling of hot material and gloves of appropriate material and thickness;
- · Clean your gloved hands with a germicide detergent;
- · Always use the special tool, supplied as standard, to extract the trays from the sterilization chamber;
- · Avoid any contact of trays and material with contaminated and/or non-heat-resistant surfaces;
- · Handle the sterile material making sure not to damage any packages, bags and containers serving as a barrier.

FOR THE PATIENT

Contamination due to unsterilised material caused by wrong cleaning treatment before sterilization.

See chapter TREATING THE MATERIAL BEFORE STERILIZATION.

Contamination due to implementation of a wrong reuse process.

Make sure to use sterile material.

Contamination due to material unsuitable to sterilization or not compliant with instructions for use.

- · Check that the contaminated material is compatible with the selected sterilization process;
- · Immediately separate the materials to be sterilized from those that must not be subjected to such process or are not able to withstand it.

Contamination due to unsterilised material caused by wrong final assessment of sterilization process.

The sterilization process electronic control system monitors the various phases, at the same time checking that the various parameters are respected; if any type of anomaly is encountered during the cycle, the program is immediately interrupted, generating an alarm identified by a code, with a relative message explaining the nature of the problem.

Furthermore, the sterilization process can be checked by means of:

CHEMICAL INDICATORS

that monitor the sterilization process by providing information, together with the control of physical and biological parameters, on the conditions occurred in the sterilization chamber during the process.

The final toning of the process indicator does not certify that the product is sterile but only that the device has been subjected to a sterilization process. If the toning does not occur, the operator in charge of releasing the sterile material, that must not be used, must find out why.

PHYSICAL INDICATORS

They include the reading of machine data and the execution of specific tests indicated during the validation phase for that specific cycle/load/autoclave. This control system can include:

- Direct reading of the synoptic system (thermometer, pressure gauge, recorder, etc.);
- Reading of prints/labels/files on which the data detected by the synoptic system are stored (parameters);
- Execution of specific tests (vacuum test, Bowie-Dick test, Helix test).

The operator in charge of the process certifies the validity of the load at the end of every cycle by means of the parametric release.

Contamination due to missing or wrong scheduled maintenance.

The sterilizer, based on a preset programming, displays a warning message relating to the scheduled maintenance necessary to ensure the good operation of the device.

Contamination due to missing periodic validation.

See chapter PERIODIC STERILIZER VALIDATION.

PACKAGE CONTENT



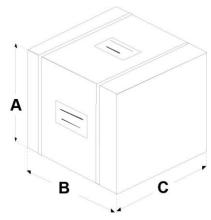
Check the integrity of the product package upon receipt.

2.1. DIMENSIONS AND WEIGHT

Once the package is opened, check that:

- The supply matches the specifications of the order (see the delivery note);
- There is no visible damage to the product.

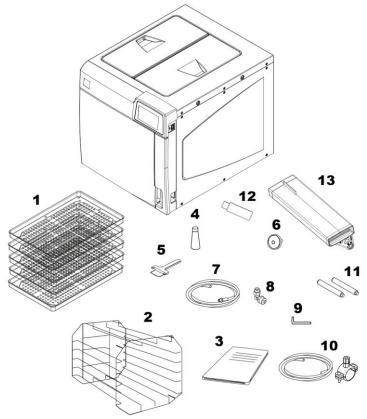
Dimensions and weight	
A Height	600 mm
B Width	600 mm
C Depth	700 mm
Total weight	68 kg





 $In \ case \ of \ wrong \ delivery, \ missing \ parts \ or \ any \ type \ of \ damage, \ inform \ immediately \ and \ in \ detail \ the \ reseller \ and \ the \ carrier \ that \ made \ the \ delivery.$

2.2. DESCRIPTION OF THE CONTENT



In addition to the sterilizer, the package contains:

- Instrument trays: 1
 - 5 pcs for 17 and 22
 - 6 pcs for 28;
- Tray holder support; 2
- 3 Operator's documentation and safety valve's EC Declaration of
- Lubricant for door locking mechanism;
- Tray extractor; 5
- Additional bacteriological filter;
- Rubber hose with quick coupling for manual water drainage; 7

- Angled union + straight union; 8
- Allen wrench (for door manual unlocking); 9
- 10 Plastic tube for direct water drainage, with fastening clamp;
- Rear spacers; 11
- USB key containing: 12 User manual and DataSter Software;
- Integrated demineralisation filter.

9

2.3. PRODUCT HANDLING

The packed product must be handled using, where possible, suitable mechanical means (lift truck, pallet truck, etc.) and following the indications on the package.

In case of manual handling, the product must be lifted by two persons using the suitable available means.

Once the sterilizer has been removed from the package, it must be lifted by two persons using the suitable available means and handled, if possible, using a truck or similar means.



We recommend to transport and store the device at a temperature not below 5°C. Extended exposure to low temperatures may damage the product.



Store the original package and use it for any transport of the device. Using a different package may damage the product during shipping.



Before transport, leave the device turned off for about 30 minutes after the last program finishes and drain the filling and discharge tanks so that all the internal parts will have time to cool down.

2.4. CONDITIONS FOR STORAGE AND TRANSPORT

TEMPERATURE: between +5° C and +70° C

HUMIDITY: between 20% and 80% **PRESSURE**: between 50 and 110 kPa

GENERAL DESCRIPTION - PRODUCT PRESENTATION

3.1. GENERAL CHARACTERISTICS

The device is an electronic water steam sterilizer, entirely operated by a micro-processor, with a large, printed stainless steel sterilization chamber. It is characterized by an advanced fractionated vacuum system for the complete removal of air, even from hollow, porous materials, and an effective final vacuum drying phase capable of eliminating all traces of humidity from any load.

The exclusive steam generation system, the effective hydraulic circuit and the electronic management (integrated by high-precision sensors) ensure a high execution speed of the process and an excellent stability of thermodynamic parameters.

Moreover, its Process Evaluation System constantly monitors all the machine's "vital" parameters in real-time, guaranteeing absolute safety and a perfect result.

The device offers users 6 sterilization programs (one of which completely programmable), all equipped with customisable, optimised drying for the fast, effective sterilization of the various types of load (instruments and materials) used in a medical environment.

All the cycles can immediately be selected on the clear LCD screen, which also allows extensive configuration of the device according to the user's needs.

For the first time, a lighting system is available for the working area in front of the sterilization chamber, which makes the daily operations easier and more comfortable.

Like in the best tradition, also the new range of autoclaves features the most complete and advanced safety systems available today, to ensure the user against any operation, electrical, mechanical, thermal or functional fault.



For the description of safety devices, refer to technical characteristics appendix.

3.2. TECHNICAL SPECIFICATIONS

3.2.1. SUMMARY TABLE

Davisa	WATER STEAM STERILIZER			
Device	17	22	28	
Class (according to Directive 93/42/EEC and subsequent amendments)		IIb		
Manufacturer		CEFLA s.c.		
Manufacturer	Via Sel	Headquarters lice Provinciale 23/A – 40026 Imola (BO) IT	
		220 V - 240 V~ 50 Hz	,	
Input voltage		220 V - 230 V~ 60 Hz		
National for a		120V~ 60 Hz		
Network fuses (6.3 x 32 mm)		F 15A 250V		
Electronic board fuses		ansformer primary 220/240 V~ 50 Hz		
(5 x 20 mm)	F2: T3.15	5A 250V (transformer primary 120 V-	~ 60 Hz)	
Nominal power		2300 W		
		1440 W (120V~ / 60 Hz)		
Insulation class		Class I		
Installation category (according to EN 61010)		Cat. II		
		Indoor use		
Operational environment	HUMID LOCATION (EN 61010 extended environmental conditions)			
A-weighted sound power level	(2140		10113)	
(ISO 3746)		< 67 db (A)		
Degree of protection (IP Code) (EN 60529:1991+A1:2000+A2:2013)	IP21			
Environmental operating conditions	Temperature: +15°C ÷ +35°C Relative humidity: between 20% and 80% max non-condensing Altitude: min -100 m / max. 3000 m (a.s.l.) Pressure: min 980 hPA / max 1045 hPA (a.s.l.)			
External dimensions (HxWxD) (rear connections excluded)	500 x 480 x 600 mm			
· · · · · · · · · · · · · · · · · · ·				
Net weight:		101		
unladen unladen, with tray holder support	approx. 48 kg approx. 51 kg	approx. 49kg approx. 52 kg	approx. 50kg approx. 53 kg	
and trays	approx. or kg	approx. 52 kg	арргох. оо ку	
unladen, with tray holder support,	50.1			
trave and water at MAN level	approx. 53 kg	approx. 654 kg	approx. 55 kg	
trays and water at MAX level	арргох. 53 кд	approx. 654 kg	approx. 55 kg	
Sterilization chamber dimensions (D	арргох. 53 kg 250 x 350 mm	approx. 654 kg 250 x 450 mm	approx. 55 kg 280 x 450 mm	
Sterilization chamber dimensions (D x D)	250 x 350 mm	250 x 450 mm	280 x 450 mm	
Sterilization chamber dimensions (D		,,,		
Sterilization chamber dimensions (D x D)	250 x 350 mm approx. 17 l (0.017 cu. m)	250 x 450 mm approx. 22 l (0.022 cu. m)	280 x 450 mm approx. 28 l (0.028 cu. m)	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support	250 x 350 mm approx. 17 l	250 x 450 mm approx. 22 l	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted)	250 x 350 mm approx. 17 l (0.017 cu. m) approx. 10 l (0.010 cu. m)	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m)	280 x 450 mm approx. 28 l (0.028 cu. m) approx. 19 l (0.019 cu. m)	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support	250 x 350 mm approx. 17 l (0.017 cu. m) approx. 10 l	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I	280 x 450 mm approx. 28 l (0.028 cu. m) approx. 19 l (0.019 cu. m)	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted) Sterilization chamber usable	250 x 350 mm approx. 17 I (0.017 cu. m) approx. 10 I (0.010 cu. m) 17 I (1.38x1.55x2.97) dm / 6.4 cu.	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m) 22 I (1.38x1.55x3.97) dm / 8.5	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I (0.019 cu. m) 28 I (1.72x1.66x3.96) dm / 11.3	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted) Sterilization chamber usable dimensions	250 x 350 mm approx. 17 I (0.017 cu. m) approx. 10 I (0.010 cu. m) 17 I (1.38x1.55x2.97) dm / 6.4 cu. dm	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m) 22 I (1.38x1.55x3.97) dm / 8.5 cu. dm approx. 5.5 I (water at MAX level)	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I (0.019 cu. m) 28 I (1.72x1.66x3.96) dm / 11.3 cu. dm	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted) Sterilization chamber usable dimensions Water tank capacity (filling)	250 x 350 mm approx. 17 I (0.017 cu. m) approx. 10 I (0.010 cu. m) 17 I (1.38x1.55x2.97) dm / 6.4 cu. dm	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m) 22 I (1.38x1.55x3.97) dm / 8.5 cu. dm approx. 5.5 I (water at MAX level) approx. 1 I (water at MIN level)	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I (0.019 cu. m) 28 I (1.72x1.66x3.96) dm / 11.3 cu. dm	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted) Sterilization chamber usable dimensions Water tank capacity (filling)	250 x 350 mm approx. 17 I (0.017 cu. m) approx. 10 I (0.010 cu. m) 17 I (1.38x1.55x2.97) dm / 6.4 cu. dm	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m) 22 I (1.38x1.55x3.97) dm / 8.5 cu. dm approx. 5.5 I (water at MAX level) approx. 1 I (water at MIN level) rd programs + 1 program defined by	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I (0.019 cu. m) 28 I (1.72x1.66x3.96) dm / 11.3 cu. dm	
Sterilization chamber dimensions (D x D) Sterilization chamber total volume Sterilization chamber usable volume (with tray holder support inserted) Sterilization chamber usable dimensions Water tank capacity (filling) Sterilization programs	250 x 350 mm approx. 17 I (0.017 cu. m) approx. 10 I (0.010 cu. m) 17 I (1.38x1.55x2.97) dm / 6.4 cu. dm	250 x 450 mm approx. 22 I (0.022 cu. m) approx. 13 I (0.013 cu. m) 22 I (1.38x1.55x3.97) dm / 8.5 cu. dm approx. 5.5 I (water at MAX level) approx. 1 I (water at MIN level) rd programs + 1 program defined by Helix/BD TEST	280 x 450 mm approx. 28 I (0.028 cu. m) approx. 19 I (0.019 cu. m) 28 I (1.72x1.66x3.96) dm / 11.3 cu. dm	

	WATER STEAM STERILIZER			
Device	17	22	28	
USB connection	Key capacity lower than or equal to 2GB: FAT formatting with 16K/sector Keys capacity higher than 2GB: FAT32 formatting with 16K/sector			
Printer connection	Serial RS232 (printer cable max length 2.5 m)			
Printer insulation class:		Class I or Class II		
Printer power supply standard:	(The safety of the sterilizer i	Compliant with Standard EN 60950. may be compromised in case of uncertif	ied printer power supply unit)	
120 V 60 Hz Main power cord	S	NEMA 5-15 plug 125 V-15A JT cable 14 AWG / 3C STYLE 1015 60 C19 connector according to IEC 60320		
220-240 V 50 Hz Main power cord	C19 conr	Plug CEE 7 / VII IEC 250V-16A 50 Hz 3x1.5 sq.mm cable from -25 to 70° C nector according to IEC 60320 UL 498, 0		
220 V 60 Hz Main power cord:		NEMA 6-15P plug 250V-15A SJT 14 AWG / 3C 300V 60° C C19 connector according to IEC 60320)	
Ethernet connection		RJ45 (max. cable length 29 m)		
Wi-Fi	802.11 b/s	g/n (2.4 Ghz); WEP / WPA / WPA2-PSK	encryption	
Bacteriological filter (filter element in PTFE)	Porosity: 0.027 microns Connection: male connector 1/8" NPT			
Maximum flow of drained water Temperature of drained water Maximum temperature of drained water	1 l/min. 50° C			
Total heat in Joule sent by the sterilizer to the surrounding air in 1 hour of continue operation	90° C 17 I = 3.6 MJ 22 I = 4 MJ 28 I = 5.4		28 I = 5.4 MJ	
Manoeuvre/handling space		1 m x 1 m		
Device	17	22	28	
Class (according to Directive 2014/68/EU PED)	Category I	Category II	Category II	
Working pressure	-0.8 ÷ 2.4 barg	-0.8 ÷ 2.4 barg	-0.8 ÷ 2.4 barg	
Safety device set	2.4 barg	2.4 barg	2.4 barg	
PT	700 kPa (abs)	700 kPa (abs)	700 kPa (abs)	
PS	2.4 barg	2.4 barg	2.4 barg	
TS	10 ÷ 140 °C	10 ÷ 140 °C	10 ÷ 140 °C	
Fluid Group	2	2	2	

3.3. SAFETY DEVICES

The sterilizer is equipped with the following safety devices for which we provide a brief description of their function:

· Mains fuses (see data in summary table)

Protection of the whole device against possible failures of heating elements.

Action: power supply interruption.

• Electronic circuit protection fuses (see data in summary table)

Protection against possible failures of the primary circuit of the transformer and of low voltage users.

Action: interruption of one or more low voltage circuits.

· Thermal circuit-breakers on mains voltage windings

Protection against possible overheating of pump motors and of transformer primary winding.

Action: temporary cut-off (until cooling) of the winding.

Safety valve

Protection against overpressure in the sterilization chamber.

Action: release of the steam and restoration of the safety pressure.

· Safety thermostat with steam generator manual reset

Protection against steam generator overheating.

Action: cut-off of the electricity to the steam generator.

· Safety thermostat with chamber heating element manual reset

Protection against overheating of the heating elements of the container under pressure.

Action: cut-off of the electricity to the chamber heating element.

· Door position safety microswitch

Confirmation of the correct closing position of the door of the container under pressure.

Action: signalling of wrong door position.

· Motor-driven door lock mechanism with electromechanical protection (pressure switch)

Protection against accidental opening of the door (even in a blackout).

Action: prevents accidental opening of the door during a program.

· Door locking mechanism safety microswitch

Striker for the correct closing position of door locking system.

Action: signalling of failed or wrong operation of door locking mechanism.

· Self-levelling hydraulic system

Plumbing system structure for the spontaneous levelling of the pressure in the case of a manual interruption of the cycle, alarm or blackout. Action: automatic restoration of atmospheric pressure in the sterilization chamber.

· Integrated system for evaluating the sterilization process

Continuous verification of the sterilization process parameters entirely managed by microprocessor.

Action: immediate interruption of the program (in case of malfunction) and generation of alarms.

· Sterilizer operation monitoring

Real-time oversight of all significant parameters when the machine is powered.

Action: generation of alarm messages (in the case of anomaly) with possible interruption of the cycle.

3.4. WATER SUPPLY CHARACTERISTICS

The sterilizer is equipped with a demineralisation filter fitted inside the filling tank, which allows supplying the device with standard water from the mains. The quality of the water treated by the integrated filter is checked automatically through a conductivity sensor.

If the demineralisation filter is not present, ONLY use demineralised/distilled water having the following characteristics to supply the sterilizer.

DESCRIPTION	VALUES IN THE WATER SUPPLY	VALUES INSIDE RESIDUAL
DRY CONDENSATE	< 10 mg/l	< 1 mg/l
SILICON OXIDE SiO ₂	< 1 mg/l	< 0.1 mg/l
IRON	< 0.2 mg/l l	< 0.1 mg/
CADMIUM	< 0.005 mg/l	< 0.005 mg/l
LEAD	< 0.05 mg/l	< 0.05 mg/l
HEAVY METAL RESIDUES (iron, cadmium	< 0.1 mg/l	< 0.1 mg/l
and lead excluded)		
CHLORIDES	< 2 mg/l	< 0.1 mg/l
PHOSPHATES	< 0.5 mg/l	< 0.1 mg/l
CONDUCTIVITY AT 20°C	< 15 μS/cm	< 3 μS/cm
pH VALUE	5 - 7	5 - 7
ASPECT	colourless, transparent, without sediment	colourless, transparent, without sediment
HARDNESS	< 0.02 mmol/l	< 0.02 mmol/l



When buying distilled water, make always sure that the quality and characteristics declared by the manufacturer are compatible with those specified in the table.



The use of water for steam generation with presence of contaminants

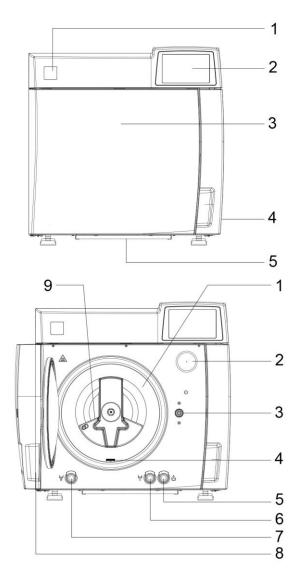
with levels exceeding those indicated in the above table can greatly shorten sterilizer lifetime.

This could also result in an increase of oxidation in the most sensitive materials as well as in an increase of limescale residues on generator, boiler, internal supports, trays and instruments.

3.5. FRONT

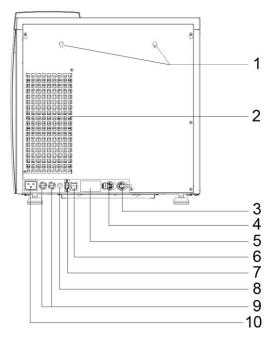
- Model
- 2 Control panel and LCD screen
- 3 Door
- 4 5 Power switch
- Dust filter

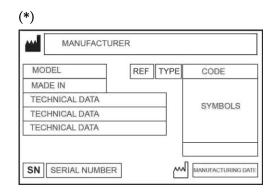
- Sterilization chamber
- Bacteriological filter 2
- Door locking system Water drainage filter 3
- 4
- 5
- Water top-up quick connector Water drainage quick connector 6
- 7 Waste water drainage quick connector
- 8 Door
- 9 Steam diffuser



3.6. **REAR**

- 1 Fastening slots for rear spacers
- 2 Heat exchanger
- 3 Connection for direct water drainage
- 4 Connection for automatic distilled water filling (only for PURE 100 / 500 AUX SV and automatic filling accessory kit)
- 5 Data plate SERIAL NUMBER LABEL (See image *)
- 6 Ethernet cable connection (max length 29 m)
- 7 Serial cable connection
- 8 Automatic filling electrical connection (only for PURE 100 / 500 AUX SV and automatic filling accessory kit)
- 9 Network fuses
- 10 Power cable connection

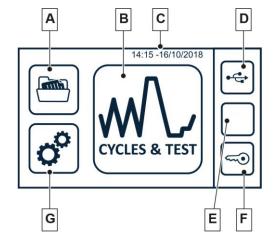




3.7. LCD ICONS

The screens in the following pictures may vary in shapes and colours, but their contents are the same as shown on the sterilizer display.

- Α Selection for data management
- Selection of sterilization and test cycles
- С Time and date
- D Button for quick download of new cycles
- E System info selection
- F Door unlock
- G Selection of the sterilizer settings (Setup)



Other particular symbols relating to the various conditions of use will be described in the relative paragraphs.

3.8. EXAMPLE OF WORKING CYCLE

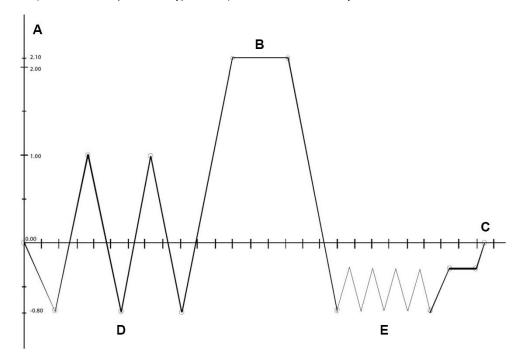
The sterilisation program can be effectively described as a succession of phases, each one with a very precise objective.

For example, the universal program (cycle B, 134°C - 4'): after loading the material in the chamber, closing the door, selecting the program and starting the cycle (after locking the door opening mechanism), the following sequence will be suggested (see the graph below):

- Preheating the generator and sterilization chamber;
- 2 Removing the air and penetration of steam in the material through a series of vacuum (extraction of the fluid from the sterilization chamber) and pressure (injection of steam into the chamber) phases;
- Raising the pressure, with the consequent increase in the temperature of the steam, until reaching the conditions required for sterilization (in the 3 example, 134°C);
- Stabilizing the pressure and temperature;
- Sterilizing for the required time (in the example, 4 minutes); 5
- Depressurizing the sterilization chamber;
- 7 Vacuum-drying phase;
- 8 Ventilating the load with sterile air;
- Bringing the pressure of the sterilization chamber back to the atmospheric level. 9

Having reached this last phase, you can unlock the door and remove the load from the sterilization chamber.

It should be emphasized that phases 1, 3, 4, 6 and 9 are identical in all cycles, with slight variations of duration that are solely dependent on the quantity and consistency of the load and the heating conditions of the sterilizer while phases 2, 5, 7 and 8 clearly vary their configuration and/or duration on the basis of the cycle selected (and, as a consequence, the type of load) and the choices made by the user.



- PRESSURE (BAR)
- В **PROCESS**
- C TIME (MIN)
- FRACTIONATED VACUUM D
- VACUUM DRYING

Please refer to the programs appendix for more details on programs.

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4. SETTING UP THE DEVICE



The safety of every system which integrates the device is responsibility of the system assembler.

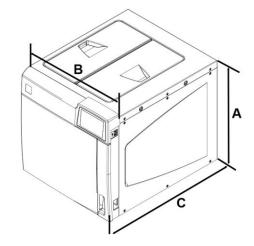
The first and essential step for a proper operation of the sterilizer, its durability over time and complete use of its features is a correct and careful commissioning. Moreover, this precaution will avoid the danger of physical injury or property damage, not to mention malfunctions and damage to the device

Please follow **meticulously** the instructions contained hereafter in this chapter.



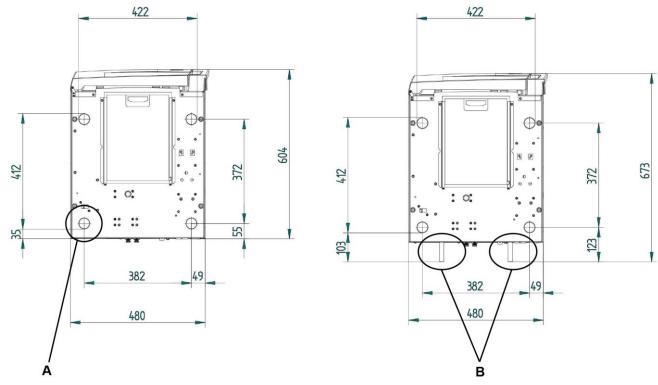
Technical service department (see appendix) is available for any doubt or further information. The sterilizer is placed on the marked only after having passed all the checks required. It does not require any additional calibration for commissioning.

Dimensions and weight		17 I	22 I	28 I
Α	Height (total)	500 mm		
В	Width (total)	480 mm		
С	Depth (excluding rear connections) Note: the sterilizer can be positioned on a surface only 550 mm deep	600 mm		
Tot	al weight	50 kg	55 kg	60 kg



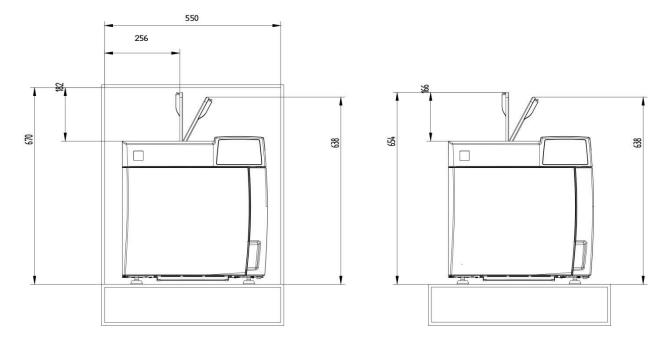
4.1. OVERALL DIMENSIONS

Centre distance and maximum overall dimensions of the sterilizer feet, with and without rear spacers.



A Feet

B Rear spacers



4.2. COMPARTMENT DIMENSIONS FOR BUILT-IN INSTALLATION

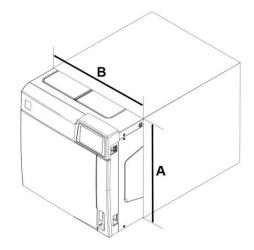
When installing the sterilizer inside a cabinet, you must provide adequate space all around the device to provide effective ventilation as well as an opening in the back (180 sq.cm) that, in addition to allowing the passage of the power cord, will also provide an adequate air flow and the consequent optimum cooling of the heat exchanger.



Mount the rear spacers supplied to ensure that the sterilizer is placed at the correct distance from the wall.

The compartment where the sterilizer will be installed must have the following minimum dimensions:

COMPARTMENT DIMENSIONS	CHAMBER VOLUME 17-22-28 L	
	520 mm	
Height	WITH FRONT FILLING OR AUTOMATIC	
	FILLING KIT	
	670 mm	
	WITH TOP FILLING	
	(FILLING DOOR OPENING)	
Width	550 mm	
Depth	600 mm	





Compartment dimensions lower than those shown may compromise the correct circulation of air around the device and may not provide adequate cooling, with the consequent deterioration of performance and/or possible damage.



If the main switch is inaccessible when installed in the compartment, use an electric plug that incorporates an on/off switch. Do not remove the upper cover nor any other external part. The device must be completely installed in the compartment. Please refer to appendix "technical characteristics" for complete technical data.

4.3. GENERAL PRECAUTIONS FOR INSTALLATION

To ensure a correct operation of the device and/or avoid risk situations, respect the following warnings:

- · Install the sterilizer on a flat and perfectly horizontal surface;
- · Make sure that the support surface is strong enough to support the device weight (about 90 kg, complete with water in hydrostatic test configuration) and has the following minimum dimensions: Width 550 mm, Depth 600 mm;
- Leave adequate space for ventilation all around the sterilizer, in particular in the rear area;
- If the device is built-in into a cabinet, be sure to respect the warnings in the previous paragraph, avoiding any obstructions of the air intakes;
- · Do not install the sterilizer too close to tubs, sinks or similar places, avoiding contact with water or liquids. This could cause short circuits and/or potentially dangerous situations for the operator:
- · Do not install the sterilizer in excessively humid or poorly ventilated environments;
- Do not install the machine in environments with flammable and/or explosive gasses or vapours;
- · Install the device so that the supply cable is not bended or squeezed.
- · It must freely run all the way to the electrical outlet;
- · Install the device so that any external filling/drainage pipes are not bent or squeezed.

4.4. POWER SUPPLY

The electrical system to which the sterilizer will be connected must be suitably dimensioned according to the electrical characteristics of the device. Plate data are shown in the TECHNICAL CHARACTERISTICS table and on the back of the machine.

4.5. ELECTRICAL CONNECTIONS

This information is shown on the back of the machine.

The sterilizer must be connected to a socket of the electric system having adequate capacity for the absorption of the device and properly earthed, in accordance with laws and/or regulations in force.

The socket must be properly protected through magneto-thermal and differential circuit breakers having the following characteristics:

Rated current I_n

Residual current I_{Dn}
 0.03 A



The manufacturer is not responsible for any damage caused by the installation of the sterilizer with unsuited and/or not properly earthed electric systems.



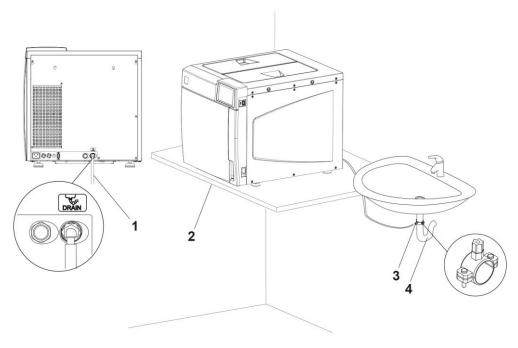
Always connect the power cord directly to the power outlet. Do not use extensions, adapters or other accessories.

4.6. DIRECT CONNECTION TO A CENTRALIZED DRAINING POINT

- · Remove the cap holding clip and the cap on the rear of the autoclave;
- Fit the plastic tube on the elbow union (supplied);
- · Fit the union and then refit the clip;
- · Fasten the clamp (supplied) to the drain siphon;
- · Cut the tube to the right length and insert its free end into the centralized draining point union locking it with the dedicated ring nut.

Make sure that the tube is not bent, crushed or obstructed in any way.

The following diagram provides an indicative arrangement of the components:



- 1 At the centralized draining point;
- Resting surface;

- 3 Clamp;
- 4 Drain siphon;



The position of the union of the centralized draining point must be lower than the resting surface of the sterilizer. Otherwise, the tank may not be emptied correctly.



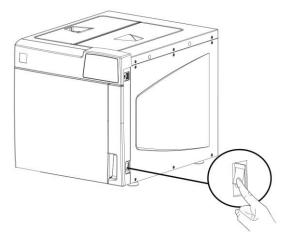
If an automatic filling system (external pump or solenoid valve, Pure 100, Pure 500) is connected the use of the direct drain connection is required. In case of fault or failure, this system allows any excess water produced by the automatic filling system to flow into the centralized draining point, thus preventing flooding.

FIRST START-UP

The time required to start the sterilizer is approximately 30 seconds.

5.1. STARTING

Once the sterilizer has correctly been installed, turn it on with the main switch on the right-hand side of the machine.





Do not turn on the sterilizer if USB key is inserted.

When the device is first turned on, the display shows the selection of LANGUAGE, DATE and TIME settings.



Once LANGUAGE, DATE and TIME have been set, the PREHEATING screen appears.

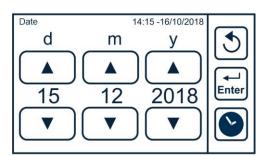
See section PREHEATING in chapter SETTINGS to set the relevant parameters.

Select the field with ▲▼ and confirm using ENTER.

Use buttons $\blacktriangle \blacktriangledown$ to adjust the values.

Confirm using ENTER and adjust the other fields.

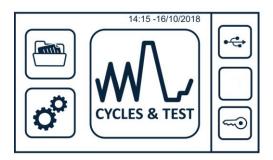




5.2. MAIN MENU

At the end of starting procedure the main menu is displayed on the side.

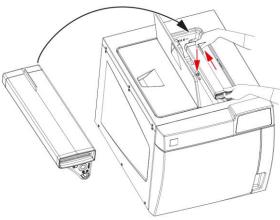
The sterilizer waits for the program selection (see "Program selection" Chapter).



5.3. INTEGRATED DEMINERALISATION FILTER

This device has been designed to demineralise drinkable water from the water mains to make it usable for the supply of the steam sterilizer. The INTEGRATED DEMINERALISATION FILTER is an ion-exchange water treatment system integrated in the autoclave's filling tank. It is a cutting-edge device with an attractive design which combines high performance, user-friendliness and a small size compared to similar water treatment systems. Its robust bacteriostatic plastic case, compliant with EN 22196 standard, and its ergonomic compact shape allow maintaining a high hygiene level in a light and small-sized filter, thanks to the installation system inside the filling water tank.

For a proper operation of the INTEGRATED DEMINERALISATION FILTER, the first essential condition is a correct installation, which avoids any malfunction or damage to the device. Please follow meticulously the instructions contained hereafter.



Only install the demineraliser on compatible autoclaves. The technical assistance service is at your disposal for further information or any doubt you may have

5.3.1. SAFETY PRECAUTIONS

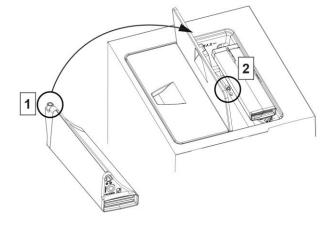
In case of contact with the ion-exchange resins inside the INTEGRATED DEMINERALISATION FILTER cartridges, follow the instructions hereafter.

CONTACT HAZARD			
Contact with eyes.	Irritating to eyes (R36).		
Contact with skin.	Lightly irritating to skin.		
FIRST-AID MEASURES			
Contact with eyes.	Rinse immediately with plenty of water and remove all particles.		
Contact with skin.	Take off contaminated clothing.		
Contact with skin.	Remove all particles and rinse affected areas with water.		
ACCIDENTAL DISPERSION MEASURES			
	Keep people away.		
recautions for people.	Pay attention to avoid falling on the slippery floor.		
Decontamination methods.	Collect the product and place it in suitable plastic containers for recycling or disposal, according to the indicated provisions.		
NOTES FOR WASTE DISPOSAL			
The used product is a special non-hazardous waste.			
The product must be disposed of according to the local, regional or national laws in force.			
The CER number for saturated or exhausted ion-exchange resins used to prepare drinkable water or water for industrial use is: 190905			

5.3.2. FIRST INSTALLATION OF THE DEMINERALISER INSIDE THE TANK

Extract the INTEGRATED DEMINERALISATION FILTER from its package, then proceed as follows:

- 1 Empty the water tank, if equipped, through the special pipe;
- 2 Insert hole (1) in the metal union (2) located at the bottom of the autoclave's tank and apply a light pressure;



- 3 Push the front part of the INTEGRATED DEMINERALISATION FILTER downwards until hearing a "clack"; this sound indicates that the filter is correctly fitted inside the tank;
- 4 Fill the tank with water from the mains;
- 5 The sterilizer is ready for use.

5.3.3. PRECAUTIONS TO AVOID WATER STAGNATION

When it is not used for long periods, the filling tank must be emptied.



Failure to empty the integrated demineralisation filter will cause the deterioration of water quality and favour bacterial growth due to water stagnation inside demineraliser's pipes and sterilizer's tanks.

TECHNICAL SPECIFICATIONS 5.3.4.

Operating temperature	+15°C ÷ +35°C
Storage temperature	+5°C ÷ +30°C
Cartridge weight	1.2 kg
Use	Internal
Inlet water	Mains drinkable water
Outlet water quality	Conductivity < 15µS/cm
Demineralised water production	Approx. 110 litres

5.4. MAINTENANCE OF THE INTEGRATED DEMINERALISATION FILTER

The integrated demineralisation filter is a consumable part, a message is displayed when it must be replaced (see alarm code table).

For replacement, see chapter DESCRIPTION OF MAINTENANCE OPERATIONS, section CLEANING AND DISINFECTION OF WATER FILTERS AND TANKS.

5.5. FILLING DISTILLED WATER

The sterilizer is equipped with a demineralisation filter fitted inside the filling tank, which allows supplying the device with standard water from the mains. The quality of the water treated by the integrated filter is checked automatically through a conductivity sensor.

If the demineralisation filter is not present, ONLY use demineralised/distilled water having the following characteristics to supply the sterilizer.

DESCRIPTION	VALUES IN THE WATER SUPPLY	VALUES INSIDE RESIDUAL
DRY CONDENSATE	< 10 mg/l	< 1 mg/l
SILICON OXIDE SiO ₂	< 1 mg/l	< 0.1 mg/l
IRON	< 0.2 mg/l l	< 0.1 mg/
CADMIUM	< 0.005 mg/l	< 0.005 mg/l
LEAD	< 0.05 mg/l	< 0.05 mg/l
HEAVY METAL RESIDUES (iron, cadmium	< 0.1 mg/l	< 0.1 mg/l
and lead excluded)		
CHLORIDES	< 2 mg/l	< 0.1 mg/l
PHOSPHATES	< 0.5 mg/l	< 0.1 mg/l
CONDUCTIVITY AT 20°C	< 15 μS/cm	< 3 μS/cm
pH VALUE	5 - 7	5 - 7
ASPECT	colourless, transparent, without sediment	colourless, transparent, without sediment
HARDNESS	< 0.02 mmol/l	< 0.02 mmol/l

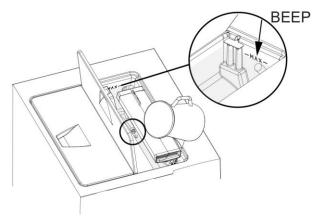
If the autoclave is to be used with distilled water, remove the integrated demineralisation filter from the filling tank.

5.5.1. **MANUAL FILLING**

The first time the sterilizer is used, and later when the lack of water is signalled, you will have to fill, or top-up, the filling water tank. Open the filling door.

Pour in water taking care not to exceed the maximum level indicated inside the tank (MAX). Close the door.

Pay attention not to spill water on the machine; in case, promptly dry.





The tank must be filled before the cycle starts or after its completion. Do not open the tank doors during the cycle execution in order to prevent hot water leaks.

AUTOMATIC FILLING 5.5.2.

Refer to the "ACCESSORIES" appendix.

6. CONFIGURATION

The sterilizers offer a wide range of customisable options. The user can thus configure the device according to his/her own needs, adapting the performance based on, for example, the type of activity carried out, the type of material to be sterilised and the frequency of use. Using the configuration program, the user can set a series of options available in user-friendly menus.



Use the configuration program whenever necessary.

A correct customisation of the device provides the best performance and the most satisfactory use.

The technical service department (see appendix) is available to help users by providing suggestions or advices on the best way to use the options in the configuration program.

6.1. SETTINGS

To enter the configuration program, select the icon shown on the side.



6.1.1. LANGUAGE

Select the LANGUAGE option.



Select the desired language scrolling the list with arrows (\blacktriangle and \blacktriangledown) and confirm by pressing ENTER.

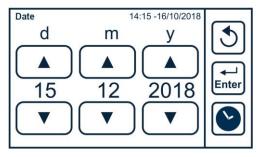


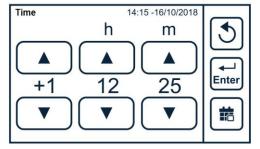
6.1.2. DATE AND TIME

Select DATE AND TIME option.

Select the field to be modified using the arrows and confirm using ENTER.







6.1.3. REMINDER

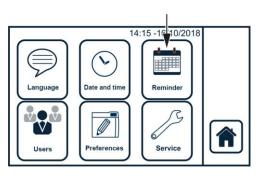
This function allows displaying, according to the selected interval, a message recalling the execution of the relative test. Select the REMINDER option.

Select if and when the Test reminders (Vacuum - Helix/B&D - Vacuum + Helix/B&D) should be enabled, using the available options. Once the fields have been set, confirm with ENTER.

Reminders activate at 8 a.m. of the selected day or upon switching-on (if it occurs after 8 a.m.).

The user can choose between:

- · Starting the test;
- Postponing the test (the reminder appears again the next day);
- Ignoring the test (the reminder appears again at the next interval).





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6.1.4. **USERS**

Press USERS button to enter the menu.

When using the device for the first time create the ADMIN user (identified by *) following the indications in the figure on the side.

Fill in the fields inserting ADMIN user name and PIN.

Press ENTER to confirm.

The first user entered is given administrator rights.



If the user enters the pin incorrectly for 3 times, it is necessary to use the unlocking procedure described in APPENDIX - USER PIN RESET.

After you have entered the PIN, you can access the reserved administrator menu.

The ADMIN user can decide whether the sterilizer will ask for the generic user PIN at the cycle start (PIN Start) and/or at the end of the cycle (PIN End).

It is possible to activate either one of the two options or both.

By activating "PIN Start" the system requests the user to enter the PIN when the sterilization cycle starts.

By activating "PIN End" the system requires to enter the PIN at the end of the cycle before unlocking the door.

If PIN request is set at cycle start, press START to select the user and the relevant PIN.

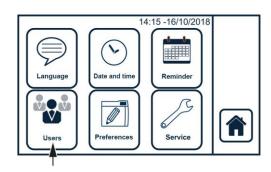
Once the PIN is confirmed, the cycle starts automatically.

To create a new user, press the "+ User" button. Fill in the fields inserting user name and PIN.

Press ENTER to confirm.

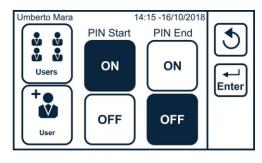
6.1.4.1. USERS LIST

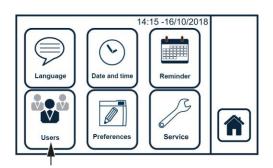
Press USERS button to enter the menu.











Select the desired user.

You will access the screen containing data of the selected user.

Once entered, a generic user can see only a summary of his/her data, or change his/her PIN (see entering PIN - the following is requested in sequence: current PIN, new PIN, new PIN confirmation).

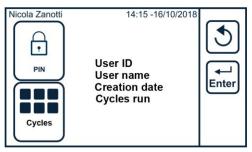
Accessing the USERS list and selecting a generic user.

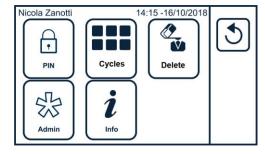
The ADMIN can instead:

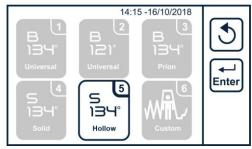
- · Give a generic user administration rights;
- Delete the user (a pop-up will be displayed to confirm the deletion);
- · View Info on users;

Select the cycles that the selected user is enabled to perform, by pressing the related icons.





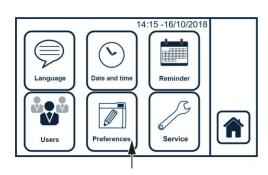


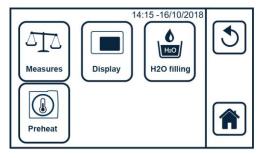


6.1.5. PREFERENCES

Press PREFERENCES button to enter the menu and set:

- UNITS OF MEASUREMENT
- DISPLAY
- WATER FILLING
- PREHEATING

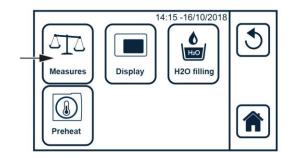




6.1.5.1. UNIT OF MEASUREMENT

Press the MEASUREMENTS icon to set the desired units of measurement (temperature, pressure), the time (12 or 24 hours) and the date format using the cursors shown in the figure.

Press ENTER to confirm the settings.





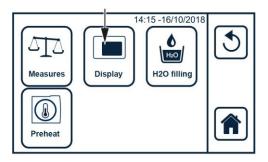
6.1.5.2. DISPLAY

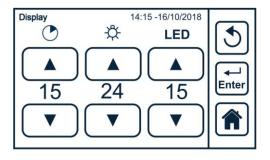
Press the DISPLAY icon to select the screen and LED bar settings. The three cursors respectively adjust:

• Screen Saver activation time-out;

- Screen brightness;
- · LED bar off time-out.

Press ENTER to confirm the settings.



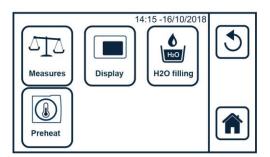


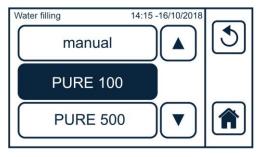
6.1.5.3. WATER FILLING

Press the H2O FILLING icon to choose the type of water filling.

Available options include:

- · Manual filling
- Pure 100
- Pure 500
- External pump
- Aux Ev kit
- · Automatic filling







When connecting the automatic filling system, the sterilizer asks you to identify the type of device actually connected by pressing the corresponding button.

If connecting the filling system when the sterilizer is off, access the menu via the configuration program and manually select the correct option.



This menu can also be used to temporarily deactivate the automatic filling system (filters exhausted, fault, etc.) and go to manual tank filling, keeping the automatic filling system connected.

6.1.5.4. PREHEATING

Select the PREHEAT. icon to activate PREHEATING in the sterilization chamber.

H₂O H2O filling Display Measures

14:15 -16/10/2018

Select ON to activate PREHEATING.

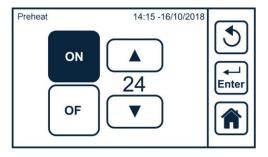
Confirm by pressing ENTER.

The arrows allow setting the maximum operating time, after which the warming up is disabled.



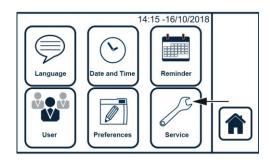
The preheating menu is displayed when the sterilizer is started for the first time, after selecting the language and date/time

The preheating is active (on) by factory default.



6.1.6. **SERVICE**

This menu is intended for the technical service department. It can be used only by an authorised technician.



7. PREPARATION OF THE MATERIAL



Always use personal protective equipment.



First of all, it should be recalled that, when handling and managing contaminated material, it is a good idea to take the following precautions:

- Wear rubber gloves of suitable thickness and the specific protective mask on your face;
- · Clean your gloved hands with a germicide detergent;
- · Always carry the instruments on a tray;
- · Never carry them in your hands;
- Protect your hands from contact with any sharp points or edges; this will avoid the risk of contracting a dangerous infection;
- Immediately remove any article that does not need to be sterilized or that is not capable of withstanding the process;
- · Carefully wash your still gloved hands when done handling non-sterile material;
- All materials and/or instruments to be sterilized must be perfectly clean, without any type of residue (deposits of organic/inorganic material, fragments of paper, cotton/gauze pads, lime, etc.).

In addition to causing problems during sterilization, the failure to clean and remove residue can damage the instruments and/or sterilizer itself.

7.1. TREATING THE MATERIAL BEFORE STERILIZATION

An effective cleaning consists of the following:

- 1 Separate metal instruments by type of material (carbon steel, stainless steel, brass, aluminium, chromium, etc.), to avoid electrolytic oxidation-reduction.
- 2 Clean the instrument with an ultrasound device containing a mixture of water and germicide solution carefully following the manufacturer's recommendations, or use a heat disinfector.
 - For best results, use a detergent specifically designed for ultrasound washing.
- 3 Manual washing is necessary if no dedicated devices are available or when automatic washing is not permitted due to the technical features of the treated material. This technique exposes the operators in charge to higher risks, for this reason it must only be applied when it is strictly necessary.
- Solutions containing phenols or quaternary ammonia compounds can cause corrosion on instruments and on the metal parts of the ultrasound device.
- 4 After washing, carefully rinse the instruments and make sure that residues have been completely eliminated; if necessary, repeat the washing cycle.
- 5 Dry all treated instruments. Drying is fundamental because the presence of water traces on the surface can jeopardise the following sterilization process.

The following items can be used for drying:

- · Paper, non-woven fabric or low-particle wipes;
- Compressed air to dry hollow instruments.

The operator must wear suitable PPE and protect the working surface to prevent its contamination by any air-dispersed particles.



To avoid the formation of lime spots, rinse with deionized or distilled water, if possible.

Whenever very hard tap water is used, we recommend always drying the instruments.

For handpieces (turbines, contra angles, etc.), in addition to the procedure described above, perform a cleaning treatment on the special devices ensuring a proper internal cleaning (sometimes including lubrication).



At the end of the sterilization program, remember to lubricate the internal handpiece mechanisms. By taking this precaution, the instrument life time will not be reduced in any way.



Consult the instructions provided by the manufacturer on the instrument/material to be sterilized before subjecting it to autoclave treatment, checking for any incompatibilities.

Strictly follow instructions for use of detergents or disinfectants and instructions for use of automatic devices for washing and/or lubrication.

As regards textile materials (porous), such as lab coats, napkins, caps and other, carefully wash and dry them before treating them in the autoclave.



Do not use detergents with a high content of chlorine and/or phosphates. Do not bleach with chlorine-based products. These substances can damage the tray supports, trays and any metal instruments that may be present in the sterilization chamber.

7.2. ARRANGING THE LOAD



Always use personal protective equipment.

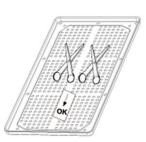




To get the best effectiveness of the sterilization process and preserve the material over time, increasing its useful life, follow the instructions below.

General notes for the positioning on trays:

- Arrange instruments made of different metals (stainless steel, hardened steel, aluminium, etc.) on different trays or anyway on trays well separated from one another.
- In case of instruments not made of stainless steel, put a sterilization paper napkin or a muslin cloth between instrument and tray, avoiding direct contact between the two different materials;
- In any case, arrange the objects sufficiently spaced from each other, so that they can remain in such position for the whole sterilization cycle;
- · Make sure that all instruments are sterilized in an open position;
- Position cutting instruments, (scissors, scalpels, etc.) so they can not come into contact with each other during sterilization; if necessary, use a cotton cloth or a gauze to isolate and protect them;
- Arrange recipients (glasses, cups, test tubes, etc.) resting on their side, or upended, thus avoiding pooling water;
- Do not load trays beyond the limit indicated (see Appendix).
- Do not stack trays one on top of the other and do not put them in direct contact with the walls of the sterilization chamber.
- · Always use the supplied tray support.
- To insert and remove trays from the sterilization chamber, always use the special supplied extractor.







Place one sterilization chemical indicator per tray to indicate when the process is complete: this will allow avoiding an unnecessary repetition of the process on the same load or, worse, the use of <u>unsterilised material</u>. If <u>packed material</u> is sterilized, place the indicator <u>inside</u> one of the packages.

Note for rubber and plastic hoses:

- · Always rinse before use with pyrogen-free water; do not dry;
- Arrange hoses on tray so that their ends are not obstructed or squashed;
- Do not bend or twist hoses, but leave them as linearly stretched as possible.



Notes for packages:

- Arrange packages next to each other, duly spaced and not stacked, avoiding their contact with chamber walls;
- Should it be necessary to wrap special objects, always use a suitable porous material (sterilization paper, muslin napkins, etc.), closing the package with adhesive tape suitable for autoclave.



Notes for packed material:

- Individually pack the instruments or, in case several instruments are placed inside the same bag, make sure they are made of the same metal;
- Seal the case with a thermosealer or adhesive tape for autoclaves;
- Do not use metal staples, needles or the like, as sterility could be affected;
- Lay the bags so as to avoid the creation of air pockets, which could potentially prevent steam correct penetration and removal;
- Position bags in such a way to leave the paper side up and the plastic side down (tray side);
- In any case, make sure that this position proves effective, reverting it, if necessary:
- If possible, using a suitable support, position bags at right angles with trav.
- · Never stack bags one on top of the other.



Always pack instruments if they have to be stored for a long time. Refer also to the indications given in chapter "sterilized material storage".

Program selection is an essential operation for the correct performance of the sterilization process.

Since all instruments, or material in general, have a different structure, consistency and properties, the **most suitable program must be identified**, both to preserve the physical characteristics (avoiding or, in any case, limiting its alterations) and to ensure the best effectiveness of the sterilization process. A guide for the selection of the correct program based on the load is present **inside Programs Appendix**.



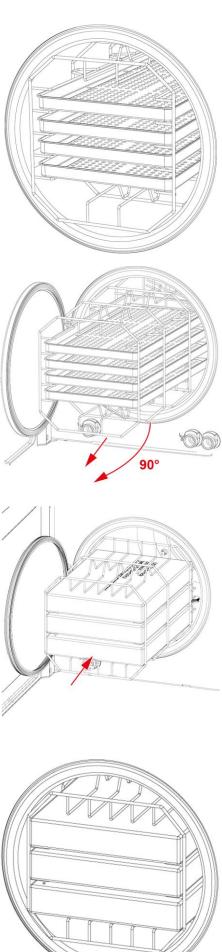
7.3. POSITIONING AND USE OF TRAY HOLDER SUPPORT

Tray holder support can be used in "tray" version (5/6 compartments based on the sterilizer model).

Or, if tray holder support is extracted and turned by 90° , it can be used to house special "boxes" (3/4 compartments based on sterilizer model).



In any case, it is possible to position the boxes (3 or 4 depending on the sterilizer model) vertically.





8. STERILIZATION CYCLES

A sterilization cycle consists of a determined number of phases.

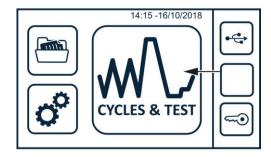
The number and duration of the phases can differ for the different cycles, based on the type of air extraction, sterilization process and drying methods:

- B 134°C Universal
- B 121°C Universal
- · S 134°C Prion
- S 134°C Hollow
- S 134°C Solid
- · Custom (User-defined)

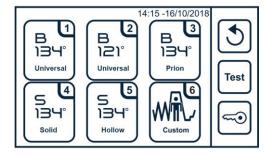
The electronic control system monitors the various phases, at the same time checking that the various parameters are respected; if any type of anomaly is encountered during the cycle, the program is immediately interrupted, generating an alarm identified by a code, with a relative message explaining the nature of the problem.

With this type of control, when you select a suitable sterilization program, you are guaranteed an effective sterilization under any conditions.

After inserting the load in the sterilization chamber (taking the precautions described in the section "**Preparing the material to be sterilized**"). Press CYCLES & TEST button to display the cycle selection buttons.



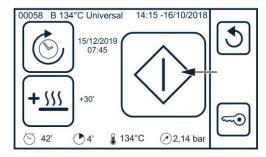
Press the button corresponding to the cycle chosen.



Start the cycle by pressing the indicated START button. The cycle counter appears in the upper left corner.

The following data are shown at the bottom:

- · Total cycle time;
- · Process time;
- · Rated process temperature.
- · Rated process pressure

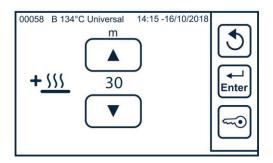


8.1. EXTRA DRYING

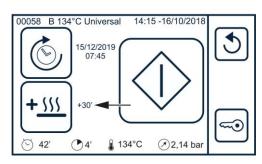
To set parameters, press and hold the following button:

00058 B 134°C Universal 14:15 -16/10/2018 07:45 +30 ~ 42' (4' 134°C 2,14 bar

Set the minutes of drying you want to add to the standard drying time and confirm with ENTER.



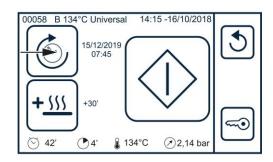
The selected value appears near the key. Activate the cycle.



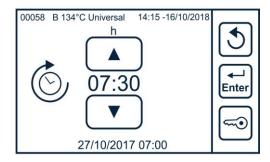
At next use, just press the extra drying button to activate the values previously set.

8.2. DELAY START

To set parameters, press and hold the following button:

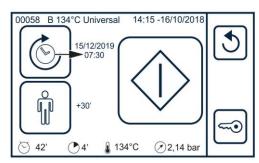


Set the time you want to start the cycle selected and confirm with ENTER.



The selected time appears near the button.

Press the START button; the cycle will automatically start at the set time.



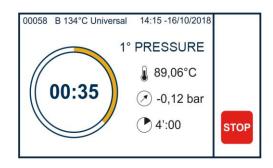
At next use, just press the delay start button to activate the values previously set.

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8.3. EXECUTION OF THE CYCLE

Taking as example the most <u>complete</u> and <u>significant</u> sterilization cycle, i.e. the **B 134°C UNIVERSAL** program, characterised by fractionated prevacuum, the cycle sequence is as follows:

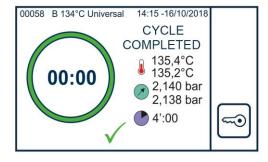
WARMING UP
FIRST VACUUM PHASE
FIRST PRESSURE RISE
SECOND VACUUM PHASE
SECOND PRESSURE RISE
THIRD VACUUM PHASE
THIRD PRESSURE RISE
STERILIZATION
STEAM DISCHARGE
DRYING
VENTILATION
CYCLE COMPLETION



8.4. CYCLE OUTCOME

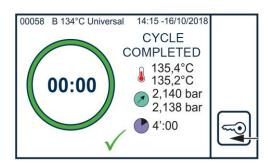
At the end of the cycle it is important to check the sterilization process outcome.

If the message "COMPLETED" is displayed, it means that the cycle has been completed correctly without any interruptions for alarms and that complete asepsis of the material is guaranteed.



8.5. DOOR OPENING AT THE END OF THE CYCLE

To open the sterilizer door, press the button shown in the figure:



8.6. USER-DEFINED CYCLE

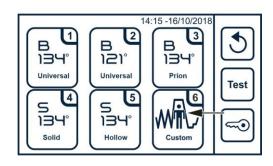
To set parameters, select the following button:

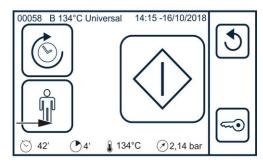
Hold down the following button to access settings:

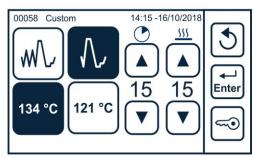
Select the type of pre-vacuum (fractionated or single), the process temperature, the exposure time and total drying time.

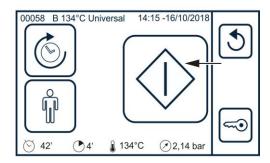
Once you have made the selections, using ENTER button, save settings and go back to the previous screen.

Press START key to start the user-defined cycle.









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9. MATERIAL STORAGE

The sterilized material must be adequately treated and stored to maintain its sterility over time, until its use.

Inadequate storage can cause rapid recontamination.

This leads to problems regardless of what you do since you will either be using recontaminated material (most of the time unconsciously), placing the user and patient at risk, or you will have to run the sterilization cycle again, with an inevitable waste of time and resources.

For this reason, we think it will be useful to provide several basic suggestions, leaving the operator the task of further study of specific texts.

Assuming that the sterilizer is located in a clean place, free of dust and not too damp, the following **precautions** should be taken when handling and/or carrying sterile material:

- 1 Remove the load from the sterilization chamber wearing gloves and a clean, or even better, sterilized smock. As an additional precaution, wear a protective mask on your face;
- 2 Rest the trays on a dry, suitably clean and disinfected surface. Take care to distance or, at any rate, separate the sterile material from the area where contaminated material is kept waiting to be sterilized;
- 3 Touch the material and/or instruments as little as possible, taking extreme care not to cut or damage the wrappings;

Let the instruments cool before any transport (and subsequent storage). If necessary for transport, transfer the material using dry, clean and disinfected containers.

The containers must be closed or, if open, covered with clean cloths.

Before use, sterile material must be stored using the appropriate techniques.

These will significantly **slow** recontamination:

- 1 Store the material and/or instruments in the protective wrappings that were used during sterilization. Do not wrap the instruments after sterilization since, in addition to being useless and completely senseless, is also potentially harmful;
- 2 Store the material in a dry, suitably clean and disinfected place, far from the area where infected material passes. If possible, use closed compartments equipped with ultraviolet light;
- 3 Identify the sterile material by attaching the sterilization date (enclosing a copy of the printed report or an adhesive label);
- 4 First use the material that has been stored the longest (FIFO, "First In First Out"). This results in material that is homogeneously stored, avoiding storing it for too long, with the consequent risks.
- 5 Never store material for too long. In fact, do not overlook the fact that materials will tend to degrade and be recontaminated in a finite time, even when the above instructions are followed.



Consult the specifications provided by the manufacturer of the packaging material relative to the maximum allowed storage time.



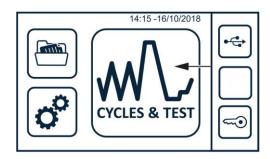
Such storage times may vary from country to country, according to the local legal requirements.

10. TEST PROGRAMS

To protect the safety of users and patients, a fundamental process like sterilizing medical devices should be periodically checked.

The device offers the possibility of easily and automatically executing two distinct test cycles:

- · HELIX TEST / B&D:
- · VACUUM TEST;
- A program that executes the two tests combined VACUUM + HELIX TEST / B/D is also available;
- · There is also a further test to check the water quality: H2O TEST.



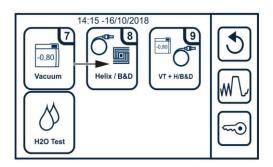
10.1. HELIX TEST/B-D CYCLE

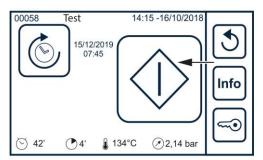
Helix/B&D Test is a cycle run at 134°C characterised by a sterilization phase that lasts a specific time (3.5 minutes); the cycle comprises the fractionated vacuum phases similar to those used in the sterilization cycles.

Using an appropriate device, you can assess correct steam penetration into hollow loads (Helix Test).

The cycle is also suitable to measure steam penetration into porous loads (Bowie & Dick test pack).

To select the **Helix/B&D Test** cycle, press the corresponding button and then press Start.

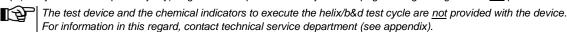




The HELIX test device (in accordance with EN 867-5 specifications) consists of a 1.5 m-long PTFE tube, with an inside diameter of 2 mm to whose end a small hermetically-sealed screw cap is fastened, able to contain an appropriate chemical indicator.

The other end of the tube is left free so that the steam can penetrate and you can assess its effectiveness.

To conduct the test (with reference to standard EN 13060), insert the chemical indicator, consisting of a paper strip with a special reagent ink in the device cap (always to be used perfectly dry). Tighten the cap in such a way that seepage through the gasket is not possible.



Place the device roughly in the middle of the central tray. Do not insert other material in the chamber. Close the door and start the cycle.

The test cycle takes place with a succession of phases similar to those described for a normal sterilization cycle.

At the end of the cycle, remove the test device from the chamber, open the cap and remove the indicator from its housing.

If the steam has correctly penetrated, the ink will have completely changed its original colour over the entire length of the strip; if not (insufficient penetration), there will only be a partial colour change or even no change at all.

The same cycle can be used for the Bowie & Dick test at the same time by placing the test device next to the HELIX test device.

Toning usually occurs from a light colour (beige, yellow, etc.) to a dark colour (blue, violet or black).

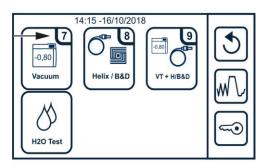
In any event, strictly follow the instructions and any additional technical details provided by the indicator manufacturer.

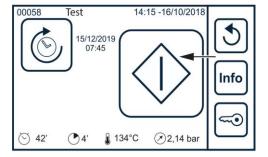
10.2. VACUUM TEST CYCLE

The VACUUM TEST cycle allows testing perfect seal of the sterilizer hydraulic system.

Measuring the variation of the degree of vacuum in a defined time-frame and comparing it with pre-established limit values, you can determine how good the seal of the sterilization chamber, tubes and the various interception devices is.

To select the VACUUM TEST cycle, select VACUUM TEST using the arrows and confirm with OK.





The cycle must be run with the sterilization chamber empty, and only the trays and their supports inserted.

We suggest to run this test at the beginning of each working day with chamber at ambient temperature.

A high chamber temperature affects the variation in the vacuum value measured during the test; the system is therefore programmed to prevent execution of the test when the operating conditions are inadequate.

Close the door and start the program.

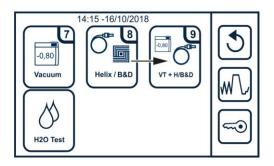
The vacuum phase starts immediately and the pressure value (bar) and the countdown from the start of the test cycle are shown on the display.

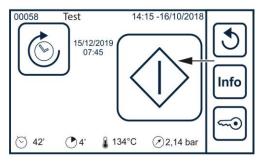
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If the pressure variation exceeds the limit defined, the program is interrupted and an alarm message generated. For the complete description of the alarms refer to appendix.

10.3. VACUUM TEST + HELIX TEST/B-D CYCLE

Select this option to run a VACUUM TEST cycle and a Helix Test/B&D cycle in sequence.





To this end, place the test device on the central tray without inserting other material.

Close the door and start the cycle.

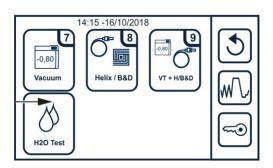
The program will execute the two cycles in succession.

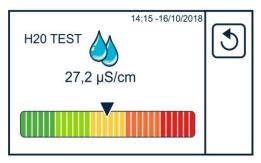
Check the results as described in the previous paragraphs.

The presence of the Helix test device and/or of the Bowie & Dick test device does not alter the execution and the result of the vacuum test cycle.

10.4. H2O TEST

Select this option to test the water quality.



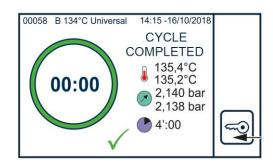


The water conductivity is automatically measured at each sterilization or test cycle start and the relative value is indicated in the cycle report.

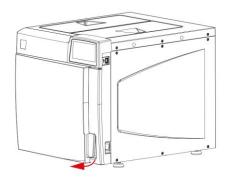
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10.5. DOOR OPENING

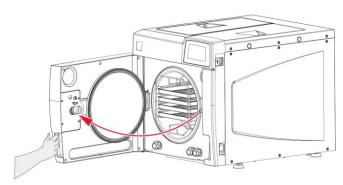
To open the autoclave door, press and hold the button shown in the figure.



The door opens and stays ajar.



Now you can manually open the door.



10.6. MANUAL INTERRUPTION

The cycle can be interrupted by the operator in any moment, by holding down the key indicated in the figure for about three seconds.



The command generates **E999 error** since the cycle could not finish correctly. Press ENTER to proceed.



If the cycle is interrupted during certain phases, an automatic cleaning procedure of the internal hydraulic circuit starts. For the complete description of the alarms refer to "alarms" appendix.

Press and hold RESET for about 3 seconds to open the door.



After a manual interruption of the program, the load must not be used since the sterilization is not ensured.

11. USED WATER DRAIN

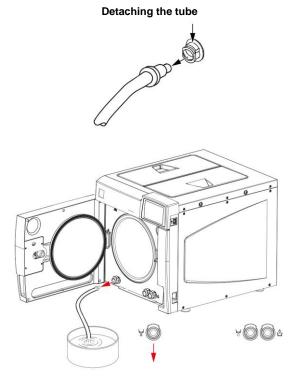
When the water maximum level is reached, a specific message is displayed.

Open the door and continue as follows:

- 1 Prepare a basin with a capacity of at least 4 litres in proximity to the sterilizer; place the free end of the drain tube provided in the basin;
- 2 Insert the other end of the tube in the female union beneath the chamber inlet (connector on the left) pushing down until you hear a click.
- 3 Completely empty out the tank and then press on the upper part of the union and detach the tube quick coupling.

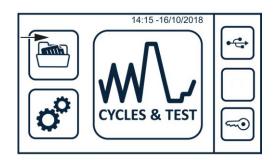


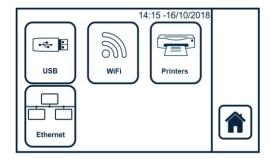
Do not open the tank doors during the cycle execution in order to prevent hot water leaks or spurts.



12. DATA MANAGEMENT

To access DATA MANAGEMENT section press the relevant icon.





12.1. USB MANAGEMENT

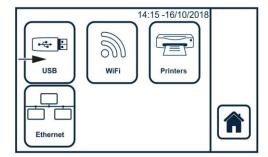
Before carrying out the following operations insert the USB key.

It is possible to copy data about the cycles carried out, stored in the inner memory of the sterilizer, on a USB key.

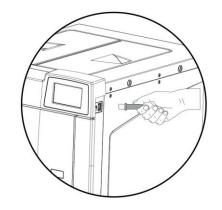
To download the files of sterilization cycles/tests select the following button:



The USB key must be formatted according to the instructions set forth in: appendix - technical characteristics, summary table.



If the USB key is not present, its insertion is required.



The sterilization / test cycle report files are in PDF format. It is possible to select the number of cycles to download on the external storage device:

- · New;
- Last 10;
- Last 50:
- Last 100;
- · Custom Mode.



If Custom Mode option is selected, you will be prompted to enter the number of the first and last cycle of the interval to download. At the end of the download it is possible to remove the key.



When a preset number of cycles is exceeded, the system generates a warning about the need to carry out a backup of data contained in the internal memory.

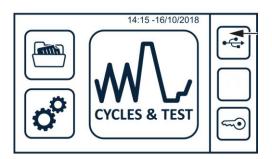
To delete the warning display, download the cycle reports using the New option.



Do not turn on the sterilizer if USB key is inserted.

12.1.1. DIRECT DOWNLOAD-NEW OPTION

To download cycles directly with NEW option, press the following button



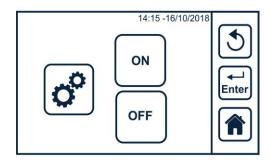
12.2. WIFI

Select Wi-Fi to connect the sterilizer to a local Wi-Fi network.

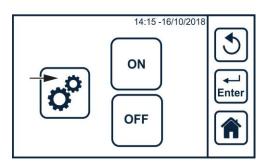
14:15 -16/10/2018
USB WiFi Printers

Ethernet

Select ON/OFF to enable or disable Wi-Fi connection.

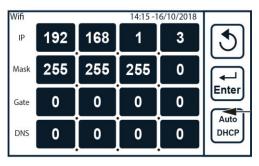


Select SETTINGS button to configure the network.



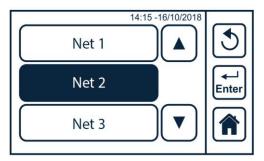
DHCP can be set as automatic or manual.

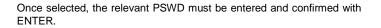
In automatic mode the network configuration parameters are assigned automatically, in manual mode the network configuration parameters must be set manually.



Choose NET SELECTION to select the network from the list automatically shown by the device.









12.3. PRINTERS

To set parameters select the following item:

Select PRINTERS to choose the support / mode to be used between:

- 1 DO NOT PRINT deactivates the printer.
- 2 REPORT allows printing a compact version of the cycle summary report at the end of the process.
- 3 EXTENDED REPORT allows printing an extended version of the cycle summary report at the end of the process.
- 4 BARCODE LABELS allows printing the labels bearing the cycle data and the barcode.

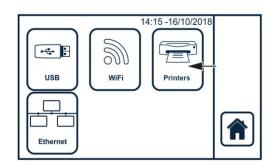
At the end of the cycle, pressing DOOR OPENING,

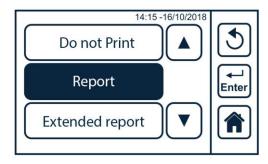
If the LABELS mode is selected, the following screen is displayed, from which it is possible to set the number of labels and the interval, in days, between the cycle execution date and the sterilised material expiry date. Use the arrows to adjust the value.

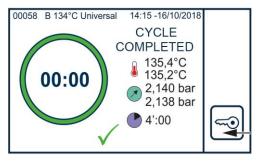
Press the PRINTER button to print the labels complete with barcode.

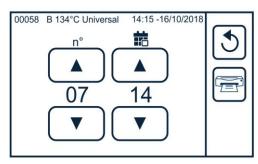
In case of unsuccessful or test cycle, only one label will be automatically printed.

If the printer is connected to the autoclave and REPORT option is set, the sterilizer automatically prints the summary report at the end of the cycle.









12.4. ETHERNET

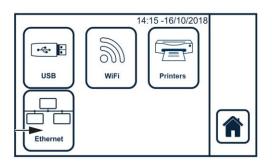
Select ETHERNET to connect the sterilizer to a local Ethernet network.

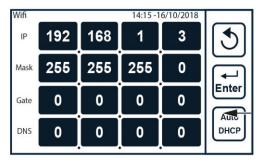
DHCP can be set as automatic or manual.

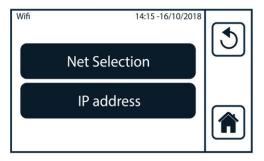
In automatic mode the network configuration parameters are assigned automatically, in manual mode the network configuration parameters must be set manually.

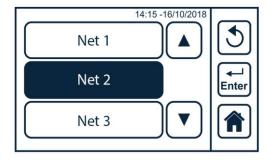
Choose NET SELECTION to select the network from the list automatically shown by the device.

Once selected, the relevant PSWD must be entered.









13. APPENDIX - PROGRAMS

Water steam sterilization is suitable for almost all the materials and instruments, provided that they can bear without damage a **minimum temperature** of 121°C (if this is not the case, other low-temperature sterilization systems must be used).

The following material can normally be sterilized with water steam:

- · Stainless steel surgical/generic instruments;
- · Carbon steel surgical/generic instruments;
- · Rotating and/or vibrating instruments driven by compressed air (turbines) or mechanical transmission (contra angles, tooth scalers);
- · Glass items;
- Mineral-based items;
- · Heat-resistant plastic items;
- · Heat-resistant rubber items;
- · Heat-resistant textiles;
- · Medication materials (gauze, pads, etc.);
- · Other generic material suitable for autoclave treatment.



Depending on the material conformation (solid, hollow or porous), on any package containing it (paper/plastic bag, paper for sterilization, container, muslin napkins, etc.) and on its resistance to heat, it is essential to choose the suitable sterilization program, referring to the table in the next page.



The device must not be used for the sterilization of fluids, liquids or pharmaceutical products.



"Prion" cycle

The reference standard for this device, EN 13060, does not lay down any requirements for inactivation processes that cause spongiform encephalopathies as scrapie, bovine spongiform encephalopathy and creutzfeldt-jakob disease.

The cycle named "prion" (18 min at 134°C) applies national regulations, which indicate this modified steam sterilization process as part of a prion decontamination program.

13.1. SUMMARY TABLE OF 17 220 V - 240 V CYCLES

	N	OMINAL	_ VALUI	ES	BAS	IC CYC	CLE P	ARAM	ETERS	STERILIZABLE N	/IATER	RIALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.00	0.30	0.30	
										Porous materials in single pack	0.75	0.25	0.25	
										Porous materials in double pack	0.60	0.20	0.20	
134°C UNIVERSAL	134	2.1	4(*)	В	F	13	40	550	0.75	Solid and hollow materials in single pack	3.00	1.00	0.50	
										Unwrapped solid and hollow materials	6.00	1.20	0.25	
										Solid and hollow instruments in double pack	1.50	0.50	0.25	
										Unwrapped porous materials	1.00	0.30	0.30	
										Porous materials in single pack	0.75	0.25	0.25	For wrapped
										Porous materials in double pack	0.60	0.20	0.20	materials and instruments (single and double pack), it
134°C PRION	134	2.1	18	В	F	13	54	600	0.85	Solid and hollow materials in single pack	3.00	1.00	0.50	is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials	6.00	1.20	0.25	
										Solid and hollow instruments in double pack	1.50	0.50	0.25	
										Unwrapped porous materials	1.00	0.30	0.30	
										Porous materials in single pack	0.75	0.25	0.25	
121°C										Porous materials in double pack	0.60	0.20	0.20	
UNIVERSAL	121	1.1	20	В	F	13	56	600	0.75	Solid and hollow materials in single pack	3.00	1.00	0.50	
										Unwrapped solid and hollow materials	6.00	1.20	0.25	
										Solid and hollow instruments in double pack	1.50	0.50	0.25	

	N	OMINA	L VALUI	ES	BAS	IC CYC	CLE P	ARAM	ETERS	STERILIZABLE I	//ATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C										Unwrapped hollow instruments	6.00	1.20	0.50	
HOLLOW UNWRAPPED	134	2.1	4(*)	S	F	4	33	550	0.65	Unwrapped solid instruments	6.00	1.20	0.50	
134°C SOLID WRAPPED	134	2.1	4(*)	S	S	13	31	350	0.55	Solid and hollow instruments "B" in single pack	3.00	1.00	0.25	It is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials "B"	6.00	1.20	0.50	
XXX°C USER (see note)	134- 121	2.1- 1.1	4÷30 - 20÷30	n.a.	F/S	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made
HELIX/BD TEST	134	2.1	3.5	-	F	1	20	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	42	-	-	-	-	-	-	

13.2. SUMMARY TABLE OF 17 120 V CYCLES

	N	OMINA	L VALU	ES	BAS	IC CY	CLE P	ARAM	ETERS	STERILIZABLE	MATE	RIALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.00	0.30	0.30	
										Porous materials in single pack	0.75	0.25	0.25	
135°C	405		4 (*)		_	40	40		0.75	Porous materials in double pack	0.60	0.20	0.20	For wrapped materials and instruments (single
HOLLOW WRAPPED	135	2.2	4(*)	В	F	13	48	550	0.75	Solid and hollow materials in single pack	3.00	1.00	0.50	and double pack), it is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials	6.00	1.20	0.25	Comiguration
										Solid and hollow instruments in double pack	1.50	0.50	0.25	
135°C SOLID UNWRAPPED	135	2.2	4(*)	s	s	4	26	350	0.55	Unwrapped solid and hollow materials "B"	6.00	1.20	0.50	
										Unwrapped porous materials	1.00	0.30	0.30	
										Porous materials in single pack	0.75	0.25	0.25	
121°C										Porous materials in double pack	0.60	0.20	0.20	For wrapped materials and instruments (single
RUBBER & PLASTIC	121	1.1	20	В	F	13	67	600	0.75	Solid and hollow materials in single pack	3.00	1.00	0.50	and double pack), it is advisable to use the 3-tray
										Unwrapped solid and hollow materials	6.00	1.20	0.25	configuration
										Solid and hollow instruments in double pack	1.50	0.50	0.25	
135° HOLLOW	405		443		_				0.5-	Unwrapped hollow instruments	6.00	1.20	0.50	
UNWRAPPED	135	2.2	4(*)	S	F	4	40	550	0.65	Unwrapped solid instruments	6.00	1.20	0.50	
135°C SOLID WRAPPED	135	2.2	4(*)	S	S	13	37	350	0.55	Solid and hollow instruments "B" in single pack	3.00	1.00	0.25	It is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials "B"	6.00	1.20	0.50	

	N	OMINA	L VALUI	ES	BAS	IC CY	CLE P	ARAM	ETERS	STERILIZABLE	MATE	RIALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
XXX°C USER (see note)	135- 121	2.2- 1.1	4÷30 - 20÷30	n.a.	F	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made
HELIX/BD TEST	135	2.2	3.5	·	F	1	24	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	50	-	-	-	-	-	-	

13.3. SUMMARY TABLE OF 22 220 V - 240 V CYCLES

	N	OMINA	L VALU	ES	BAS	IC CYC	CLE P	ARAM	ETERS	STERILIZABLE	MATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.20	0.40	0.30	
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	
134°C UNIVERSAL	134	2.1	4(*)	В	F	15	44	700	0.8	Solid and hollow materials in single pack	4.00	1.25	0.50	
										Unwrapped solid and hollow materials	7.50	1.20	0.25	
										Solid and hollow instruments in double pack	2.00	0.60	0.25	
										Unwrapped porous materials	1.20	0.40	0.30	
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	For wrapped materials and
134°C PRION	134	2.1	18	В	F	15	58	750	0.9	Solid and hollow materials in single pack	4.00	1.25	0.50	instruments (single and double pack), it is advisable to use the 3-tray
										Unwrapped solid and hollow materials	7.50	1.20	0.25	configuration
										Solid and hollow instruments in double pack	2.00	0.60	0.25	
										Unwrapped porous materials	1.20	0.40	0.30	
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	
121°C UNIVERSAL	121	1.1	20	В	F	15	61	750	0.8	Solid and hollow materials in single pack	4.00	1.25	0.50	
										Unwrapped solid and hollow materials	7.50	1.20	0.25	
										Solid and hollow instruments in double pack	2.00	0.60	0.25	

	N	OMINA	L VALU	ES	BAS	IC CYC	CLE P	ARAM	IETERS	STERILIZABLE	MATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C										Unwrapped hollow instruments	7.50	1.50	0.50	
HOLLOW UNWRAPPED	134	2.1	4(*)	S	F	5	37	750	0.7	Unwrapped solid instruments	7.50	1.50	0.50	
134°C SOLID WRAPPED	134	2.1	4(*)	S	s	15	37	400	0.6	Solid and hollow instruments "B" in single pack	4.00	1.00	0.25	It is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials "B"	7.50	1.20	0.50	
XXX°C USER (see note)	134- 121	2.1- 1.1	4÷30 - 20÷30	n.a.	F/S	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made
HELIX/BD TEST	134	2.1	3.5	-	F	1	24	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	46	-	-	-	-	-	-	

13.4. SUMMARY TABLE OF 22 120 V CYCLES

	N	OMINA	L VALU	ES	BAS	IC CY	CLE P	ARAM	ETERS	STERILIZABLE	MATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.20	0.40	0.30	
										Porous materials in single pack	1.00	0.30	0.25	
40500										Porous materials in double pack	0.75	0.25	0.20	For wrapped materials and
135°C HOLLOW WRAPPED	135	2.2	4(*)	В	F	15	53	550	0.8	Solid and hollow materials in single pack	4.00	1.25	0.50	instruments (single and double pack), it is advisable to use the 3-tray
										Unwrapped solid and hollow materials	7.50	1.20	0.25	configuration
										Solid and hollow instruments in double pack	2.00	0.60	0.25	
135°C SOLID UNWRAPPED	135	2.2	4(*)	S	s	5	32	400	0.6	Unwrapped solid and hollow materials "B"	7.50	1.50	0.50	
										Unwrapped porous materials	1.20	0.40	0.30	
										Porous materials in single pack	1.00	0.30	0.25	
121°C										Porous materials in double pack	0.75	0.25	0.20	For wrapped materials and instruments (single
RUBBER & PLASTIC	121	1.1	20	В	F	15	73	750	0.8	Solid and hollow materials in single pack	4.00	1.25	0.50	and double pack), it is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials	7.50	1.20	0.25	configuration
										Solid and hollow instruments in double pack	2.00	0.60	0.25	
135° HOLLOW										Unwrapped hollow instruments	7.50	1.50	0.50	
UNWRAPPED	135	2.2	4(*)	S	F	5	44	750	0.7	Unwrapped solid instruments	7.50	1.50	0.50	
135°C SOLID WRAPPED	135	2.2	4(*)	S	S	15	44	400	0.6	Solid and hollow instruments "B" in single pack	4.00	1.00	0.25	It is advisable to use the 3-tray configuration
WINAI I'ED										Unwrapped solid and hollow materials "B"	7.50	1.20	0.50	
XXX°C USER (see note)	135- 121	2.2- 1.1	4÷30 - 20÷30	n.a.	F	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made

	N	OMINA	L VALU	ES	BAS	IC CYC	CLE P	ARAM	IETERS	STERILIZABLE I	MATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
HELIX/BD TEST	135	2.2	3.5	-	F	1	24	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	50	-	-	-	-	-	-	

13.5. SUMMARY TABLE OF 28 220 V - 240 V CYCLES

	N	OMINA	L VALU	ES				YCLE TERS		STERILIZABLE N	/IATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.50	0.50	0.50	
										Porous materials in single pack	1.25	0.35	0.35	
134°C										Porous materials in double pack	0.90	0.30	0.30	
UNIVERSAL	134	2.1	4(*)	В	F	17	54	900	0.8	Solid and hollow materials in single pack	5.00	1.50	0.75	
										Unwrapped solid and hollow materials	9.00	1.40	0.25	
										Solid and hollow instruments in double pack	2.50	0.70	0.25	
										Unwrapped porous materials	1.50	0.50	0.50	
										Porous materials in single pack	1.25	0.35	0.35	
										Porous materials in double pack	0.90	0.30	0.30	For wrapped
134°C PRION	134	2.1	18	В	F	17	68	950	1	Solid and hollow materials in single pack	5.00	1.50	0.75	materials and instruments (single and double pack), it
										Unwrapped solid and hollow materials	9.00	1.40	0.25	is advisable to use the 3-tray configuration
										Solid and hollow instruments in double pack	2.50	0.70	0.25	
										Unwrapped porous materials	1.50	0.50	0.50	
										Porous materials in single pack	1.25	0.35	0.35	
										Porous materials in double pack	0.90	0.30	0.30	
121°C UNIVERSAL	121	1.1	20	В	F	17	67	950	0.9	Solid and hollow materials in single pack	5.00	1.50	0.75	
										Unwrapped solid and hollow materials	9.00	1.40	0.25	
										Solid and hollow instruments in double pack	2.50	0.70	0.25	

	N	OMINA	L VALU	ES				YCLE		STERILIZABLE N	/IATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWM/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C										Unwrapped hollow instruments	9.00	1.50	0.50	
HOLLOW UNWRAPPED	134	2.1	4(*)	S	F	6	42	950	0.8	Unwrapped solid instruments	9.00	1.50	0.50	
134°C SOLID WRAPPED	134	2.1	4(*)	S	s	17	43	500	0.7	Solid and hollow instruments "B" in single pack	5.00	1.00	0.25	It is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials "B"	9.00	1.20	0.50	
XXX°C USER (see note)	134- 121	2.1- 1.1	4÷30 - 20÷30	n.a.	F/S	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made
HELIX/BD TEST	134	2.1	3.5	-	F	1	24	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	46	-	-	-	-	-	-	

13.6. SUMMARY TABLE OF 28 120 V CYCLES

	N	OMINA	L VALU	ES				YCLE		STERILIZABLE N	//ATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped porous materials	1.50	0.50	0.50	
										Porous materials in single pack	1.25	0.35	0.35	
135°C										Porous materials in double pack	0.90	0.30	0.30	For wrapped materials and instruments (single
HOLLOW WRAPPED	135	2.2	4(*)	В	F	17	65	900	0.8	Solid and hollow materials in single pack	5.00	1.50	0.75	and double pack), it is advisable to use the 3-tray
										Unwrapped solid and hollow materials	9.00	1.40	0.25	configuration
										Solid and hollow instruments in double pack	2.50	0.70	0.25	
135°C SOLID UNWRAPPED	135	2.2	4(*)	S	S	6	38	500	0.6	Unwrapped solid and hollow materials "B"	9.00	1.50	0.50	
										Unwrapped porous materials	1.50	0.50	0.50	
										Porous materials in single pack	1.25	0.35	0.35	
121°C										Porous materials in double pack	0.90	0.30	0.30	For wrapped materials and instruments (single
RUBBER & PLASTIC	121	1.1	20	В	F	17	80	950	0.8	Solid and hollow materials in single pack	5.00	1.50	0.75	and double pack), it is advisable to use the 3-tray
										Unwrapped solid and hollow materials	9.00	1.40	0.25	configuration
										Solid and hollow instruments in double pack	2.50	0.70	0.25	
135° HOLLOW										Unwrapped hollow instruments	9.00	1.50	0.50	
UNWRAPPED	135	2.2	4(*)	S	F	6	50	950	0.7	Unwrapped solid instruments	9.00	1.50	0.50	
135°C SOLID WRAPPED	135	2.2	4(*)	S	S	17	52	500	0.6	Solid and hollow instruments "B" in single pack	5.00	1.00	0.25	It is advisable to use the 3-tray configuration
										Unwrapped solid and hollow materials "B"	9.00	1.20	0.50	
XXX°C USER (see note)	135- 121	2.2- 1.1	4÷30 - 20÷30	n.a.	F	5÷30	n.a.	n.a.	n.a.	Unwrapped solid instruments (other load types are possible depending on the user settings)	n.a.	n.a.	n.a.	Variable parameters depending on the settings made

	N	OMINA	L VALU	ES				YCLE TERS		STERILIZABLE N	IATER	IALS		NOTES
CYCLE DESCRIPTION	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Load Max)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	ТҮРЕ	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
HELIX/BD TEST	135	2.2	3.5	-	F	1	24	-	-	Test device only (without another load)	-	-	-	
VACUUM TEST	-	-0.8	-	-	-	-	18	-	-	Empty chamber	-	-	-	
VACUUM + HELIX/BD TEST (executable in sequence)	-	-	-	-	-	-	50	-	-	-	-	-	-	



(*) To set a sterilization time of 5.5 minutes, contact the Technical Service.

Single Pre-Vacuum = 1 pre-vacuum; -0.8 bar (see figures in the following pages).

Fractionated Pre-Vacuum = 3 pre-vacuum; -0.8 bar each (see figures in the following pages).

Definition of hollow loads in accordance with standard EN13060:2014.

The term "hallow loads" in this manual refers both to "narrow lumen" elements (paragraph 3.18 EN 13060:2014) and "simple hollow" elements (paragraph 3.30 EN 13060:2014).

The term "hollow load B" refers ONLY to the elements defined as "simple hollow" (paragraph 3.30 EN 13060:2014).

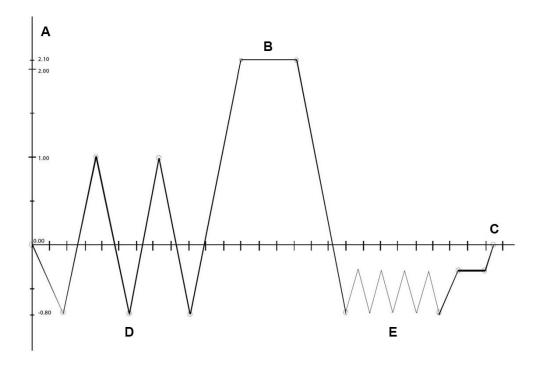
			RE, TIME AND TEMP th EN 13060: 2014 fo			
			134°C cycles			
EN 13060:2014		Time (minutes)	Min temperature	Max temperature	Min pressure (bar)	Max pressure (bar)
1	CS					
t1	1PV				-0.81	-0.79
t2	1PP				+0.97	+1.03
t3	2PV				-0.81	-0.79
t4	2PP				+0.97	+1.03
t5	3PV				-0.81	-0.79
t6	SS	4 / 5.5	+134	+138	+2.04	+2.40
t7	SE	4 / 5.5	+134	+138	+2.04	+2.40
t8	DS				-0.81	-0.79
t9	DE					
2	CE				-0.02	+0.02
<u>.</u>			121°C cycles			
EN 13060:2014		Time (minutes)	Min temperature	Max temperature	Min pressure (bar)	Max pressure (bar)
1	CS					
t1	1PV				-0.81	-0.79
t2	1PP				+0.97	+1.03
t3	2PV				-0.81	-0.79
t4	2PP				+0.97	+1.03
t5	3PV				-0.81	-0.79
t6	SS	20	+121	+125	+1.05	+1.31
t7	SE	20	+121	+125	+1.05	+1.31
t8	DS				-0.81	-0.79
t9	DE					
2	CE				-0.02	+0.02

13.7. STERILIZATION PROGRAM DIAGRAM

PROGRAM 134°C UNIVERSAL 134°C – 4' 00"

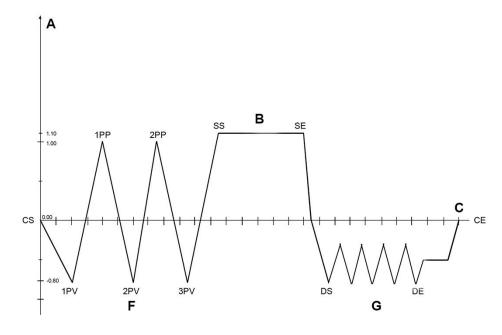
PROGRAM 134°C PRION 134°C – 18' 00"

- A PRESSURE (BAR)
- **B** PROCESS
- C TIME (MIN)
- **D** FRACTIONATED VACUUM
- **E** VACUUM DRYING



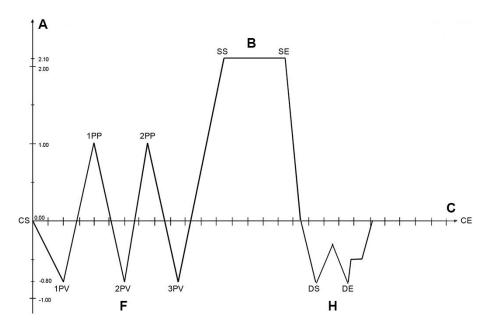
PROGRAM 121°C UNIVERSAL 121°C – 20' 00"

- A PRESSURE (BAR)
- B PROCESS
- C TIME (MIN)
- F FRACTIONATED PRE-VACUUM
- **G** LONG DRYING



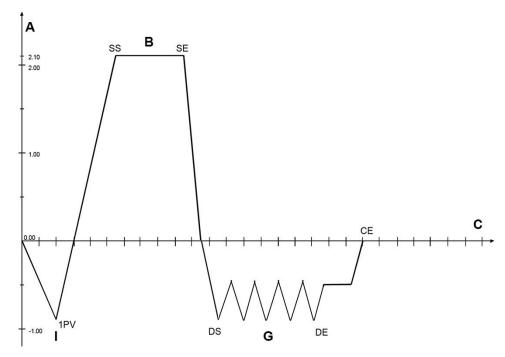
PROGRAM 134°C HOLLOW UNWRAPPED 134°C – 4'00"

- A PRESSURE (BAR)B PROCESS
- C TIME (MIN)
- F FRACTIONATED PRE-VACUUM
- H SHORT DRYING



PROGRAM 134°C SOLID WRAPPED 134°C – 4'00"

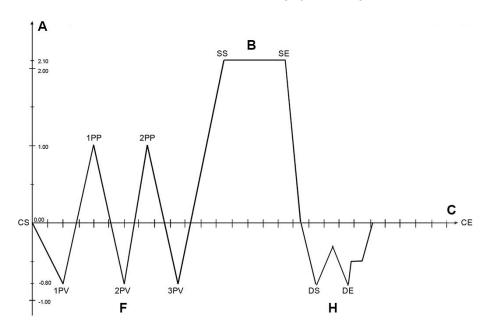
- A PRESSURE (BAR)
- **B** PROCESS
- C TIME (MIN)
- I SINGLE PRE-VACUUM
- **G** LONG DRYING



13.8. DIAGRAMS OF THE TEST PROGRAMMES

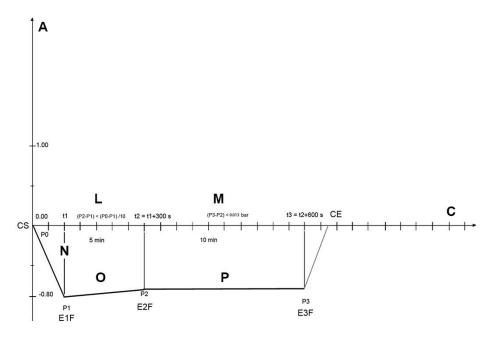
PROGRAM HELIX B&D TEST 134°C – 3'0"

- A PRESSURE (BAR)
- **B** PROCESS
- C TIME (MIN)
- F FRACTIONATED PRE-VACUUM
- H SHORT DRYING



PROGRAM VACUUM TEST -0.80 bar

- A PRESSURE (BAR)
- C TIME (MIN)
- L INTERMEDIATE CONDITION TO CONTINUE THE TEST
- M FINAL CONDITION TO PASS THE TEST
- N VACUUM PHASE
- **O** STANDBY
- P LOSS MEASUREMENT



13.9. EXAMPLES OF PRINTED REPORTS

(WITH OPTIONAL PRINTER)

PROGRAM PRINTING (STANDARD)

Model S/N Ver. SW Counter Selection Temperature Pressure Process tim Stand-by Pre-vacuum Drying	е	0007/0015 134 °C SOL 134 °C 2.10 bar 4 min LOW SINGLE FAST	.ID
CYCLE STA	RT	01/02/11 12:14	
Time		С	bar
00:01 02:02 05:48 06:02 07:02 08:02 09:02 10:02 10:37 11:41 16:08 17:12	CS 1PV ET SS SE DS SPD DE CE	079.4 093.7 135.6 135.9 135.6 135.5 135.4 135.5 104.1 047.5 047.6 084.6	+0.00 -0.80 +2.15 +2.17 +2.14 +2.14 +2.15 +0.00 -0.90 -0.84 -0.04
06:32 09:59	MAX MIN	136.0 135.4	
Drying Puls CYCLE ENI		01 01/02/11 12:36	
STERILIZA	TION:	POSITIVE	

Model	
S/N	
Ver. SW	
Counter	0007/0015
Selection	134 °C UNIVERSAL
Temperature	134 °C
Pressure	2.10 bar
Process time	4 min
Stand-by	HIGH

Stand-by Pre-vacuum Drying				
CYCLE S	TART	01/02/10 09:52		
Time		С	bar	
00:01 01:57 04:53 07:00 09:15 11:22 15:04 15:19 17:18 18:19 19:53 20:57 26:55 29:15 29:43	CS 1PV 1PP 2PV 2PP 3PV ET SS SE DS SPD EPD DE CE	075.1 047.8 120.5 061.1 120.4 061.1 135.5 135.9 135.4 135.5 135.4 135.5 104.4 094.9 112.6 115.8	-0.00 -0.80 +1.00 -0.80 +0.98 -0.80 +2.15 +2.17 +2.14 +2.15 +0.00 -0.90 -0.86 -0.47 -0.04	
25.43	CE	115.6	-0.04	

Drying Pulses CYCLE END	05 01/02/11 10:28
STERILIZATION:	POSITIVE

MAX MIN 135.9 135.4

PROGRAM PRINTING HELIX/BD TEST

Model S/N Ver. SW Counter Selection Temperati Pressure Process ti CYCLE S	me	0011/0019 HELIX TE: 134 °C 2.10 bar 3.5 min 01/02/11 16:38	
Time		С	bar
00:01 02:06 04:35 05:45 07:02 08:15 11:00 11:14 13:14 14:14 14:45 15:20 16:34 18:21 19:21 20:06	CS 1PV 1PP 2PV 2PP 3PV	076.4 089.3 120.4 062.5 120.2 061.1 135.6 136.0 135.6 135.5 135.4 111.5 047.8 059.5	+0.00 -0.89 +0.99 -0.78 +0.97 -0.79 +2.15 +2.17 +2.14 +2.14 +0.00 -0.89 -0.86 -0.50
12:33 14:44 Drying pu CYCLE E		136.0 135.4 01 01/02/11 17:01	

HELIX TEST COMPLETE
Please attach the indicator hereunder

PROGRAM PRINTING VACUUM TEST

Model S/N Ver. SW Counter Selection	S/N Ver. SW Counter		9 TEST
CYCLE S	TART	01/02/11 11:37	
Time		С	bar
00:00	cs	035.0	+0.00
01:39	E1F	037.4	-0.80
6:39	E2F	038.4	-0.79
22:39	E3F	042.0	-0.79
23:54	CE	045.5	-0.01
CYCLE E	ND	01/02/11 12:01	
VACUUM	TEST:	POSITIVE	

P

16:20 18:11

The text printed on the receipt remains for a couple of years, if correctly stored (away from any source of heat and in a cool, dry place). Store thermal receipts using cellulose envelopes, do not use plastic bags.

14. APPENDIX - MAINTENANCE

In addition to correct use, the user needs to perform ordinary maintenance in order to guarantee safe, efficient operation over the device's entire life.



Always use personal protective equipment.



For better quality of maintenance, supplement routine checks with regular periodic check-ups that can be performed by Technical Service Department (see Appendix).

It is also fundamental to perform a <u>periodic sterilizer validation</u>, i.e. a check of process thermo-dynamic parameters and their comparison with the reference values detected by duly calibrated tools. Refer to 'Sterilizer periodic validation' in the next part of the Appendix.

The ordinary maintenance described below consists in easy manual operations and preventive interventions involving simple instruments.



In the event of replacement of components or parts of the device, request and/or use original spare parts only.

14.1. ORDINARY MAINTENANCE PROGRAMME

The table summarizes the maintenance interventions required to maintain the sterilizer in good working order.

In case of **heavy use**, we recommend to **shorten** maintenance intervals:

DAILY	Clean the gasket and the internal part of the door Clean external surfaces
WEEKLY	Clean the sterilization chamber and its accessories Disinfect external surfaces Clean/disinfect filling/drainage tanks
PERIODICALLY	See Scheduled Maintenance messages
YEARLY	Validate sterilizer (see scheduled maintenance)

14.2. SCHEDULED MAINTENANCE MESSAGES

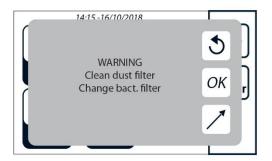
The sterilizer periodically displays warning messages relevant to "routine" maintenance operations that must be carried out in order to ensure the proper operation of the device.

Press OK to confirm that the required maintenance operation has been completed.



button to postpone the operation.

In this case, the warning message will reappear the next time the sterilizer is used.



Warnings are displayed with the following frequency:

WARNING MESSAGE

BOILER FILTER CLEANING DOOR LOCK LUBRICATION **DUST FILTER CLEANING** BACTERIOLOGICAL FILTER REPLACEMENT WATER TANK CLEANING **BOILER GASKET REPLACEMENT** GENERAL SERVICE



A regular maintenance is essential to achieve the best performance of the device.

Periodically, a message will be displayed requesting that the above maintenance operations are performed.

For further information or in case of doubt, contact the technical service: if they have performed regular maintenance on the device, the technician might have already carried out some of these operations (e.g. >Replacement of the bacteriological filter or of the gasket).

Always keep in mind the following general warnings:

- Do not wash the sterilizer with direct jets of water, neither under pressure nor sprinkled. Seepage into electrical and electronic components could damage the functioning of the device or its internal parts, even irreparably;
- Do not use abrasive cloths, metal brushes (or other aggressive materials) or products for metal cleaning, both solid and liquid, to clean the device or the sterilization chamber;
- Do not use unsuitable chemical products or disinfectants to clean the sterilization chamber. In fact, these products can cause damages, even irreparably;
- Do not allow limescale or residues of other substances to accumulate in the sterilization chamber, on the door and on the gasket, but remove them periodically. In fact, such residues may damage these parts, besides compromising the operation of the hydraulic circuit components.



The formation of white spots on the base of the internal wall of the chamber means that you are using poor quality demineralised water.



Before performing ordinary maintenance, make sure that the power cord plug is removed from the mains socket.

If this is not possible, move the external switch of the device's power supply line to Off.

If the external switch is distant or not visible to the maintainer, place a "work in progress" sign on the switch, after turning it off.



14.3. DESCRIPTION OF MAINTENANCE INTERVENTIONS

Let's now look at the various operations to be carried out.

14.3.1. CLEAN GASKET AND PORTHOLE

To eliminate any traces of limestone, clean the gasket of the chamber and the door porthole with a clean cotton cloth soaked in a soft solution of water and vinegar (or a similar product, checking the contents on the label before using).

Dry the surfaces and remove any residues before using the device.

14.3.2. CLEAN STERILIZATION CHAMBER AND ACCESSORIES

Clean the sterilization chamber, support and trays (and internal surfaces in general) with a clean cotton cloth soaked in water and, possibly, the addition of a small amount of neutral detergent.

Carefully rinse with distilled water, taking care not to leave any type of residue in the chamber or on accessories.



Do not use pointed or sharp tools to remove scale from the sterilization chamber.

Should there be evident deposits, immediately check the quality of the distilled water used (see technical characteristics appendix).

14.3.3. EXTERNAL SURFACE CLEANING AND DISINFECTION

To clean and disinfect the external surfaces, we recommend using STER 1 PLUS or ethyl alcohol diluted with 50% water. Apply product with a soaked cloth, then dry.

As an alternative, we recommend using products containing the following at no more than the given concentration:

- Ethanol. Concentration: maximum 30 g per 100 g of disinfectant.
- 1-Propanol (n-propanol, propyl alcohol, n-propyl alcohol). Concentration: maximum 20 g per 100 g of disinfectant.
- Combination of ethanol and propanol. Concentration: the combination of the two must be maximum 40 g per 100 g of disinfectant.



Do not spray or vaporise any product directly on device surfaces.

Inflammable liquid.

14.3.4. BOILER FILTER CLEANING

With use it is likely that various residues accumulate in the filter and with time obstruct the lower drain duct.

To clean the filter, open the sterilizer door and remove the cap using a coin or another suitable tool.

Loosen the union that contains the filter.

Remove the filter from its support and thoroughly clean it under a jet of running water, if necessary using a sharp tool to remove any large foreign bodies (if possible use a jet of compressed air).

If it is impossible to recover the filter, replace it with a new one.

Refit everything operating in reverse order and making sure to screw the union in such a way that the drain holes are positioned at the level of the boiler wall.



Properly fit the filter in its housing.

A partial fitting may damage the component.

14.3.5. DOOR LOCK LUBRICATION

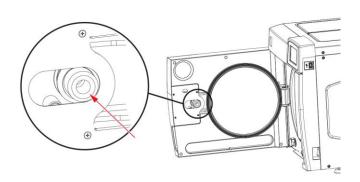
Using a clean cloth, remove any residues from the bushing and the screw. Lubricate the inside of the bushing on the sterilizer door with a film of the silicon-based grease provided (as shown in the figure).



Wear single-use gloves before application.

Essentially, the lubricant is not irritant to the skin; nevertheless, it may cause unpleasant effects if it accidentally comes into contact with eyes.

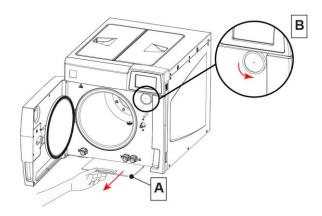
In case of contact with eyes, rinse with plenty of water.



14.3.6. DUST FILTER CLEANING

Remove the dust filter (A) from the lower part of the autoclave, thoroughly rinse it with water and dry it before refitting it.

The filter can be cleaned using a jet of compressed air, making sure not to disperse any dust into the environment.



14.3.7. REPLACE THE BACTERIOLOGICAL FILTER

When filter maintenance is due or every time you notice visible clogging of the filter (indicated by the filter markedly turning grey), unscrew the bacteriological filter (B) from its support and replace it with a new one, screwing it fully down on the union.



A spare bacteriological filter is provided with the device.

If you need spare parts of this component, refer to technical assistance appendix.

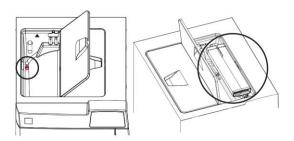
14.3.8. CLEANING AND DISINFECTION OF FILTER, INTEGRATED DEMINERALISATION FILTER AND WATER TANKS

Clean and disinfect filters and internal walls only of the tank with a disposable wipe/cloth soaked in ethyl alcohol 70%.



Do not use 70% alcohol to disinfect the other plastic surfaces.

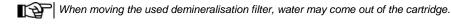
Empty the tanks containing the autoclave filling and drainage water, remove any deposits around the filters at the bottom of the tanks (see figure) and on the INTEGRATED DEMINERALISATION FILTER (if equipped). After removing and cleaning the filters, wipe the inside of the tanks with a cloth soaked in ethyl alcohol 70% and clean thoroughly. After cleaning the filters, refit them inside the tanks.



14.3.9. REPLACEMENT OF THE INTEGRATED DEMINERALISATION FILTER

When the special alarms indicate the production of demineralised water with conductivity levels above the allowed threshold (20 microsiemens), the ionexchange resin cartridge must be replaced:

- · Turn the sterilizer off;
- · Remove the upper cover;
- · Empty the filling tank of the autoclave;
- Remove the cartridge by lifting the rear part, then the front part;
- Replace the cartridge according to the initial installation procedure (see chapter INTEGRATED DEMINERALISATION FILTER).



14.3.10. BOILER GASKET REPLACEMENT

It is advisable to have the boiler gasket replaced by an authorised technician, therefore contact Technical Service (see APPENDIX - TECHNICAL SERVICE).

14.4. PERIODIC STERILIZER VALIDATION

As happens with all devices, it is possible, and sometimes inevitable, to have a decrease in performance and the effectiveness of components along their lifespan, in a period of time dependent on its frequency of use.

To guarantee the safety of the process over time, it is periodically (possibly annually) necessary to verify, the thermodynamic process parameters (pressure and temperature), to check if they continue to remain within allowed limits or not.

The requalification of the sterilizer's performance is the **responsibility of the user** of the product.

The reference European standards EN 17665 (Sterilization of the medical devices - Method for the validation and systematic control of the steam sterilization) and EN 556 (Sterilization of the medical devices - Requirements for the medical devices marked with "STERILE" indication) supply an effective guide tool for carrying out the verifications on the steam sterilizers.

Since, in addition to specific experience and training, these controls require the use of special equipment (high-precision sensors and probes, data loggers, dedicated software, etc.) suitably verified and calibrated, it is necessary to contact a company specializing in these activities.



Customer support department (see Appendix) is available to provide any information relative to the periodic validation of steam sterilizers.

14.5. DEVICE USEFUL LIFE

Water steam sterilizer service life is of 10 years (average use: 5 cycles/day, for 220 days/year). For normal use, it is expected that the device is used and maintained according to the instructions provided by the manufacturer.

The expected useful life of the device is subject to risk analysis carried out in compliance with requirements of standard ISO 14971:2012.

14.6. DISPOSING THE EQUIPMENT WHEN NO LONGER USED

According to Directive 2012/19/EU concerning waste disposal, the units must not be disposed of as municipal waste, but must be separated. When purchasing a new device of an equivalent type, one for one, the device that has come to the end of its lifetime should be returned to the dealer for disposal. As regards reuse, recycling and other forms of recovery of the above mentioned waste, the manufacturer carries out the functions defined in the individual national legislations.

Appropriate differentiated waste collection for subsequent recycling treatment and environmentally friendly disposal contributes to preventing possible negative effects on the environment and health and encourages recycling of the materials of which the device is made up. The symbol indicating separate collection for electrical and electronic equipment consists of the crossed out bin marked on the device.



Under national legislation, fines can be imposed if the product is disposed in an illegal manner.

15. APPENDIX - GENERAL PROBLEMS

If while using the device a problem or an alarm occurs, this DOES NOT mean that the device is out of order.

It may not, in fact, be related to a breakdown but, more probably to an anomalous situation, often merely transitory (such as a blackout), or incorrect use.

In any case, it is important to first identify the cause of the failure and then take suitable corrective actions, either autonomously or with the intervention of the Technical Service Department (see Appendix).

For this purpose, below, we provide instructions for diagnosing and resolving general problems, in addition to a precise description of the alarm codes, their meaning and their solution.

15.1. TROUBLESHOOTING

If your sterilizer is not working correctly, please make the following checks before contacting the Technical Service Department:

PROBLEM	POSSIBLE CAUSE	SUGGESTED SOLUTION
	The power cable is not plugged-in.	Plug it in.
	Lack of voltage at the power supply socket.	Check the cause of the lack of voltage at socket and fix it.
The sterilizer does not power-on.	The main switch and/or differential switch are turned to OFF.	Turn the switch to ON.
	The mains fuses are blown.	Replace with good fuses of equal nominal value. (See the Summary Table in Appendix, Technical Characteristics).
After pressing START, the sterilization cycle does not start.	The device is preheating.	Wait for the sterilizer to reach the proper operating conditions for starting the program. **Under standard conditions, the Average Preheating Time is about 10-15 Minutes.**
The safety valve has triggered.	Locking ring loosened. Presence of anomalous overpressure in the chamber.	Check the proper tightening of the milled ring nut of the safety valve. Let the device cool or use gloves to prevent burns while touching the valve.
Water presence on the sterilizer	The water automatic filling system hose (optional) is not correctly connected.	Check the tightness of the fittings and, if necessary, reassemble them more carefully. Check that the hoses are completely inserted on the fittings; check the presence of hose clamps.
resting surface.	Steam leak from door gasket.	At the end of the cycle clean the gasket and the closing porthole with a dampened cloth. Check the presence of any gasket damage. Perform a new verification cycle.
	Excessive load in the sterilization chamber.	Check that the load does not exceed the maximum values allowed (See the Summary Table in Appendix "Technical Characteristics").
Excessive humidity on the material and/or instruments at the end of the program.	Load not correctly positioned.	Position the load, in particular the wrapped one, as per the indications. (See <u>Chapter</u> "Preparing the material").
	Wrong selection of the sterilization program.	Choose the sterilization program suitable for the type of material to be treated. (See the Summary Table in "Programs" Appendix).
	Clogged chamber drainage filter.	Clean or replace the drainage filter. (See Appendix "Maintenance").

PROBLEM	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Quality of the instruments not adequate.	Check the quality of instruments, making sure that the material they are made of is suitable to tolerate the steam sterilization.
Traces of oxidation or spots on instruments	Organic or inorganic residues on the instruments.	Carefully clean the material before subjecting it to the sterilization cycle. (See Chapter "Preparing the material").
	Contact between instruments made of different metals.	Separate instruments made of different metals. (See Chapter "Preparing the material").
	Presence of limescale residues on the wall of the chamber and/or accessories.	Clean the chamber and the accessories as prescribed. (See Appendix "Maintenance").
Blackening of the instruments or damage to the material.	Wrong selection of the sterilization program.	Choose the sterilization program suitable for the type of material to be treated. (See the Summary Table in "Programs" Appendix).

16. APPENDIX - ALARMS



If the problem persists, contact the technical service (see APPENDIX) communicating the sterilizer model and serial number.

These data are indicated on the registration plate on the rear side of the device and on the declaration of conformity and can be viewed also by means of the "sterilizer information" command.

Every time an anomalous condition occurs during the operation of the sterilizer, an alarm is generated, identified by a specific code (consisting of a letter followed by a 3-digit number).

Alarm codes are divided into four categories:

E= ERROR/WARNING

Incorrect handling and/or use or a cause outside the device.

The problem can normally be solved by the user.

Code format: (xxx = identification number 000 ÷ 999)

A = ALARM

First level fault

The problem can normally be solved on site by a specialised technician. Code format: Axxx (xxx = identification number 000 ÷ 999)

H = HAZARD

Second level fault

The problem can normally be solved by the Technical Service Centre. (xxx = identification number 000 ÷ 999) Code format: Hxxx

S = SYSTEM ERROR

Electronic system error (HW-FW).

Code format: Sxxx (xxx = identification number 000 ÷ 999)



In case of alarm, switch off the device only after having followed the indications displayed and having carried out the reset (see "Resetting the system" paragraph).

16.1. ALARM INTERVENTION

The alarm intervention causes the cycle interruption (or the normal operation interruption), the display of the relevant alarm code and message and an audible warning.

16.2. ALARM DURING A CYCLE

The alarm procedure is designed in order not to give the user any possibility to confuse an anomalous cycle with an efficiently carried out one, and therefore to unintentionally use not sterilised materials; it is structured to guide the user up to the RESET of the sterilizer and the following use.

16.3. SYSTEM RESET

The system can be reset in two <u>alternative ways</u>, depending on the type of alarm occurred (see the List of alarm codes below in this appendix):

- Pressing the OK button;
- Following the instructions displayed and holding down the RESET button for about 3 seconds.

Press the RESET button for approx. 3 seconds to go back to the main



After the RESET and any technical operation necessary to eliminate the fault, the device will be ready to perform a new program.



Never turn off the device before carrying out the reset.

17. ALARM CODES

The $\underline{\textbf{list}}$ of alarm codes, the relevant messages displayed and RESET modes, are indicated in the following table:

17.1. ERRORS (CATEGORY E)

The alarm codes in the list can refer to functions that are not present on the models concerned in this Manual

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
		POWER OUTAGE	
E000	Black-out	CONTACT TECH.SERVICE	2
E001	Voltage of power supply line too high	OVERVOLTAGE CONTACT TECH.SERVICE	1
E002	Water conductivity threshold 1 exceeded	INSUFFICIENT H2O QUALITY	1
E003	Water conductivity threshold 2 exceeded	QUALITY H2O BAD CHANGE WATER	1
E004	Error in electrical mains frequency reading	LINE FREQ. ERROR CONTACT TECH. SERVICE	1
E007	One of the two fans is not working properly	FAN PROBLEM CONTACT TECH.SERVICE	1
E008	Water conductivity threshold 1 exceeded	FILTERS NEARLY EXHAUSTED	1
E009	Water conductivity threshold 2 exceeded	QUALITY H2O BAD CHANGE FILTERS CHANGE WATER	1
E010	Door open	DOOR OPEN CLOSE DOOR	1
E011	Water conductivity threshold 2 exceeded by filling tank	QUALITY H2O BAD CHANGE FILTERS CHANGE WATER	2
E012	The cycle threshold for the periodic replacement of integrated water filters has been reached	CHANGING FILTERS	1
E013	Water conductivity threshold 2 exceeded by discharge tank	QUALITY H2O BAD CHANGE FILTERS CHANGE WATER	2
E020	Door lock system (closing) activation time-out exceeded	DOOR CLOSING ERROR CONTACT TECH.SERVICE	1 (then reattempt or turn off)
E021	Door lock system (opening) activation time-out exceeded	DOOR OPENING ERROR CONTACT TECH.SERVICE	1 (then reattempt or turn off)
E022	Door lock microswitches failure.	DOOR LOCK PROBLEM CONTACT TECH.SERVICE	2
E030	The water in the feed tank is at minimum level (MIN)	LOAD TANK MINIMUM LEVEL FILL TANK	1
E031	Maximum level of water in the drainage tank (MAX)	DISCHARGE TANK MAXIMUM LEVEL EMPTY TANK	1
E042	The MAX water level in the filling tank has been reached	LOAD TANK MAXIMUM LEVEL	1
E050	Reminder to run the Vacuum Test cycle	TEST REMINDER RUN VACUUM TEST	1
E051	Reminder to run the Helix Test cycle	TEST REMINDER RUN HELIX TEST	1
E052	Reminder to run the combined Vacuum + Helix Test cycle	TEST REMINDER RUN VACUUM+HELIX TEST	1
E060	The autoclave cannot connect to Lan network	ETHERNET CONFIG. ERROR CHECK SETTINGS	1
-	•	•	

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
E061	The autoclave cannot connect to Wi-Fi network	WI-FI CONFIG. ERROR CHECK SETTINGS	1
E900	Vacuum test failed (during TEST PHASE)	TEST FAILED SECOND STEP CONTACT TECH.SERVICE	2
E901	Vacuum test failed (during STAND-BY PHASE)	TEST FAILED FIRST STEP CONTACT TECH.SERVICE	2
E902	Vacuum test failed (vacuum pulse time-out exceeded)	TEST FAILED VACUUM NOT ACHIEVED CONTACT TECH.SERVICE	2
E998	Remote maintenance activity in progress	REMOTE SERVICE ACTIVE	1
E999	Manually interrupting the cycle	MANUAL INTERRUPTION	2

^{1 =} OK (warning) 2 = OK + door unlocking + RESET (if in cycle)

17.2. ALARMS (CATEGORY A)

A032		FILL.WATER LEVEL	
	Problem with the level sensor of the filling tank	SENSOR PROBLEM CONTACT TECH.SERVICE	1
A033	Problem with the level sensor of the discharge tank	DISCH.WATER LEVEL SENSOR PROBLEM CONTACT TECH.SERVICE	1
A034	Filter in the water filling tank removed during cycle	FILLING TANK FILTER REMOVED DURING CYCLE	2
A035	Filter in the water drain tank removed during cycle	DRAIN TANK FILTER REMOVED DURING CYCLE	2
A040	The tank has not been filled (only with automatic filling system)	FAILED WATER INLET CHECK AUTOMATIC LOAD	1
A042	The MAX water level in the filling tank has been reached abnormally (automatic filling)	WATER FILLING MAXIMUM LEVEL CHECK TANK	1
A043	Discharge tank full beyond maximum limit	DISCHARGE TANK MAXIMUM LEVEL CHECK TANK	1
A101	Temperature sensor PT1 broken (sterilization chamber)	CHAMBER PROBE PT1 OPEN CIRCUIT CONTACT TECH.SERVICE	1
A102	Temperature sensor PT2 broken (steam generator)	GENERATOR PROBE PT2 OPEN CIRCUIT CONTACT TECH.SERVICE	1
A103	Temperature sensor PT3 broken (heating element)	HEATING BAND PROBE PT3 OPEN CIRCUIT CONTACT TECH.SERVICE	1
A105	Temperature sensor PT5 broken (conductivity measurement compensation)	CONDUCTIVITY SENSOR PT5 OPEN CIRCUIT CONTACT TECH.SERVICE	1
A111	Temperature sensor PT1 short-circuited (sterilization chamber)	CHAMBER PROBE PT1 SHORT-CIRCUIT CONTACT TECH.SERVICE	1
A112	Temperature sensor PT2 short-circuited (steam generator)	GENERATOR PROBE PT2 SHORT-CIRCUIT CONTACT TECH.SERVICE	1
A113	Temperature sensor PT3 short-circuited (heating element)	HEATING BAND PROBE PT3 SHORT-CIRCUIT CONTACT TECH.SERVICE	1
A115	Temperature sensor PT5 short-circuited (conductivity measurement compensation)	CONDUCTIVITY SENSOR PT5 SHORT-CIRCUIT CONTACT TECH.SERVICE	1
A116	ADC error	PROCESS BOARD ERROR CONTACT TECH.SERVICE	1
A120	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH.SERVICE	1
A121	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH.SERVICE	1
A122	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH.SERVICE	1
A123	Solenoid valves control driver fault	EV DRIVER ERROR CONTACT TECH.SERVICE	2
A124	Door motor control driver fault	DOOR MOTOR DRIVER ERROR CONTACT TECH.SERVICE	2
A125	Faulty current draw detected	FAULTY CURRENT DRAW CONTACT TECH.SERVICE	2
A126	Connection error with Wi-Fi module	WI-FI MODULE ERROR CONTACT TECH.SERVICE	1
A127	Communication error between graphical interface and process board via Can	CAN ERROR CONTACT TECH.SERVICE	2
A128	Communication error between graphical interface and ethernet module	ETHERNET MODULE ERROR CONTACT TECH.SERVICE	1
A129	Connection error with NFC module	NFC MODULE ERROR CONTACT TECH.SERVICE	1
		RGB LED BAR ERROR	+

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
A131	Solenoid valve 1 failed	SOLENOID VALVE 1 ERROR	2
		CONTACT TECH.SERVICE SOLENOID VALVE 2 ERROR	
A132	Solenoid valve 2 failed	CONTACT TECH.SERVICE	2
A133	Solenoid valve 3 failed	SOLENOID VALVE 3 ERROR CONTACT TECH.SERVICE	2
		SOLENOID VALVE 4 ERROR	
A134	Solenoid valve 4 failed	CONTACT TECH.SERVICE	
A135	Solenoid valve 5 failed	SOLENOID VALVE 5 ERROR CONTACT TECH.SERVICE	2
A136	Solenoid valve 6 failed	SOLENOID VALVE 6 ERROR CONTACT TECH.SERVICE	2
A137	Solenoid valve 7 failed	SOLENOID VALVE 7 ERROR CONTACT TECH.SERVICE	2
		STEAM GENERATOR	
A201	Pre-heating not executed within time-out (steam generator)	RESISTOR OPEN CIRCUIT	2
		CONTACT TECH.SERVICE	
4000		HEATING BAND	0
A202	Pre-heating not executed within time-out (tube bundle heating element)	OPEN CIRCUIT CONTACT TECH.SERVICE	2
		1PV TIMEOUT	
A250	1st pulse with vacuum not achieved within time out	CHECK LOAD	2
	·	CHECK CHAMBER FILTER	
A251	1st rise back up to atmospheric pressure not reached within the time-out	ATM1 UPSTROKE TIMEOUT	2
		CONTACT TECH.SERVICE	
A252	1st pressure pulse not reached within the time-out	1PP UPSTROKE TIMEOUT CONTACT TECH.SERVICE	2
		2PV TIMEOUT	
A253	2nd pulse with vacuum not achieved within time out	CHECK LOAD	2
	·	CHECK CHAMBER FILTER	
A254	2nd rise back up to atmospheric pressure not reached within the time-out	ATM2 UPSTROKE TIMEOUT	2
		CONTACT TECH.SERVICE 2PP UPSTROKE TIMEOUT	
A255	2nd pressure pulse not reached within the time-out	CONTACT TECH.SERVICE	2
		3PV TIMEOUT	
A256	3rd pulse with vacuum not achieved within time-out	CHECK LOAD	2
		CHECK CHAMBER FILTER	
A257	3rd rise back up to atmospheric pressure not reached within the time-out	ATM3 UPSTROKE TIMEOUT CONTACT TECH.SERVICE	2
A258	3rd pressure pulse not reached within the time out	3PP UPSTROKE TIMEOUT	2
71200	ora prosoure pulse not readined within the time out	CONTACT TECH.SERVICE	
A260	Chamber depressurization not reached within time out	ATM3 DOWNSTROKE TIMEOUT	2
A200	Chamber depressurization not reached within time out	CHECK LOAD CHECK CHAMBER FILTER	2
		PRESSURE LEVELLING	
A261	Chamber levelling not reached within time out	TIMEOUT	2
		CONTACT TECH.SERVICE	
A262	Vacuum pulgation during druing not avacuted within time out	PD PRESSURE	2
A262	Vacuum pulsation during drying not executed within time-out	UPSTROKE TIMEOUT CONTACT TECH.SERVICE	2
		ATM1 DOWNSTROKE TIMEOUT	
A353	1st drop to atmospheric pressure not completed within the time-out	CHECK LOAD	2
		CHECK CHAMBER FILTER	
A250	2nd drop to observe here are not consulated within the time.	ATM2 DOWNSTROKE TIMEOUT	
A356	2nd drop to atmospheric pressure not completed within the time-out	CHECK LOAD CHECK CHAMBER FILTER	2
		SPD PRESSURE	
A360	Vacuum pulsation after maintenance step not executed within time-out	DOWNSTROKE TIMEOUT	2
	·	CONTACT TECH.SERVICE	
		PD PRESSURE	_
A362	Chamber depressurisation during drying not reached within time-out	DOWNSTROKE TIMEOUT	2
		CONTACT TECH.SERVICE	

^{1 =} OK (warning) 2 = OK + door unlocking + RESET

17.3. HAZARDS (CATEGORY H)

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
H150	MPX pressure sensor broken/not connected	PRESSURE SENSOR OPEN CIRCUIT CONTACT TECH.SERVICE	3
H160	MPX pressure sensor short-circuited	PRESSURE SENSOR SHORT-CIRCUIT CONTACT TECH.SERVICE	3
H400	Pconv/T ratio not balanced (Pconv>T) (STERILIZATION phase)	INCORRECT P/T RATIO CHECK LOAD	2
H401	T/Pconv ratio not balanced (T>Pconv) (STERILIZATION phase)	INCORRECT T/P RATIO CHECK LOAD	2
H402	Temperature over the MAX limit (STERILIZATION phase)	TEMPERATURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	2
H403	Temperature below the MIN limit (STERILIZATION phase)	TEMPERATURE BELOW MINIMUM LIMIT CONTACT TECH. SERVICE	2
H404	Floating temperature over the limit (STERILIZATION phase)	ERRATIC TEMPERATURE CONTACT TECH. SERVICE	2
H405	Pressure over the MAX limit (STERILIZATION phase)	PRESSURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	2
H406	Pressure below MIN limit (STERILIZATION phase)	PRESSURE BELOW MINIMUM LIMIT CONTACT TECH. SERVICE	2
H410	Time measurement error	INTERNAL TIMER ERROR CONTACT TECH.SERVICE	2
H990	Excessive pressure (sterilization chamber, MPX)	PRESSURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	2
H991	Overheating (sterilization chamber, PT1)	PT1 OVERHEATING CHECK LOAD	2
H992	Overheating (steam generator, PT2)	PT2 OVERHEATING CONTACT TECH.SERVICE	2
H993	Overheating (layer resistance, PT3)	PT3 OVERHEATING CONTACT TECH.SERVICE	2

^{1 =} OK (warning) 2 = OK + door unlocking + RESET 3 = Cycle failed + OK + door unlocking + RESET

17.4. SYSTEM ERRORS (CATEGORY S)

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
		FLASH MEMORY	
S001	Flash memory 1 on process board failed	NOT ACCESSIBLE	2
		CONTACT TECH.SERVICE	
		FLASH MEMORY	
S002	Flash memory 2 on process board failed	NOT ACCESSIBLE	2
0002	radii memery 2 on process sound ranou	CONTACT TECH.SERVICE	_
		FLASH MEMORY	
S003	Flach momery 1 on graphical interface heard	NOT ACCESSIBLE	2
3003	Flash memory 1 on graphical interface board		2
		CONTACT TECH.SERVICE	
		FLASH MEMORY	_
S004	Flash memory 2 on graphical interface board	NOT ACCESSIBLE	2
		CONTACT TECH.SERVICE	
S005	LICD key not apposible	PROBLEM WITH USB KEY	2
3003	USB key not accessible	CHANGE KEY	2
		USB KEY	
S006	USB key not accessible	NOT ACCESSIBLE	2
0000	oca no, not accession	CHANGE KEY	_
		USB KEY FULL	
S007	USB key full	CHANGE KEY	2
S009	Printer not connected	PRINTER DISCONNECTED	2
		CHECK CONNECTION	
S010	Printer: there is no paper or there might be a configuration error	PRINTER PAPER OUT	2
0010	Times: there is no paper of there might be a comigaration ener	CHECK PAPER	
S011	Printer cover apen	PRINTER:	2
3011	Printer cover open	DOOR OPEN	2
0040	B 1 11 11 11 11 11 11 11 11 11 11 11 11	PRINTER: NOT READY	
S012	Probable printer configuration error	TRY AGAIN	2
		RUN BACKUP	
S020	Cycle backup not done	DOWNLOAD NEW CYCLES	2
		CYCLE MEMORY FULL	
S021	Cycle storage limit exceeded	START OVERWRITING	2
	Charle using a watchday that one of main tasks is not in aroth	SYSTEM ERROR	2 (eff evale)
S030	Check, using a watchdog, that one of main tasks is not in crash		2 (off-cycle)
	condition	CONTACT TECH.SERVICE	3 (in cycle)
S031	Check, using a hardware watchdog, that one peripheral is not in	SYSTEM ERROR	2 (off-cycle)
	lock condition.	CONTACT TECH.SERVICE	3 (in cycle)
S032	Check, using a watchdog, that one of main tasks is not in lock	SYSTEM ERROR	2 (off-cycle)
3032	condition (e.g. infinite loop)	CONTACT TECH.SERVICE	3 (in cycle)
0004	OM made mating	SYSTEM ERROR	0
S034	SW malfunction	CONTACT TECH.SERVICE	2
		SYSTEM ERROR	_
S035	SW malfunction in solenoid valve management	CONTACT TECH.SERVICE	2
		SYSTEM ERROR	2 (off-cycle)
S040	Check the log saving in the Flash memory	CONTACT TECH.SERVICE	3 (in cycle)
			3 (III cycle)
S041	Cycle performed with 4-minute sterilization time at 134°C	LOG SAVE ERROR	1
ļ	, ,	CONTACT TECH.SERVICE	
S042	Cycle performed with standard drying	4-MINUTE STERILIZATION	1
30.2	5,5.5 performed man ordinated drying	COMPLETED	'
1		STANDARD DRYING	
S099	Error during cycle report creation	CHECK	1
1		LOAD DRYING	
		PROBLEM IN CREATING	
S100	SW malfunction	CYCLE REPORT	2
3100	OV manufolion	CONTACT TECH.SERVICE	
		CONTACT TECHLOLINGE	1

^{1 =} OK (warning) 2 = OK + door unlocking + RESET 3 = Cycle failed + OK + door unlocking + RESET

17.5. TROUBLESHOOTING

According to the type of alarm occurred, please find below the indications to detect the possible causes and restore the proper operation:

The alarm codes in the list can refer to functions that are not present on the models concerned in this Manual.

17.5.1. ERRORS (CATEGORY E)

•

CODE **POSSIBLE CAUSE** SUGGESTED SOLUTION Sudden power failure (blackout). Wait for the power to be restored and do a RESET following the instructions. The main switch has accidentally been turned off Reconnect the plug and/or turn the device on again and RESET according to and/or the power plug pulled from the socket. the instructions. E000 Replace with good fuses of equal nominal value. (See the Summary Table in Network fuses blown. Appendix Technical Characteristics). Turn the device on again and RESET according to the instructions. Reset according to the instructions. If the problem occurs again, have the E001 Abnormal voltage peak on the mains. mains electric system checked by a technician. RESET according to the instructions. Empty the filling tank and refill it with E002 The filling tank contains water of inadequate quality. distilled water of adequate quality (<15µs/cm). RESET according to the instructions. IMMEDIATELY empty the filling tank and refill it with distilled water of adequate quality (<15µs/cm). In these conditions, the sterilizer allows starting a maximum of 5 E003 The filling tank contains water of very poor quality. cycles, after which it locks until the tank is filled with distilled water of adequate quality (<15 µs/cm). This precaution is necessary to prevent damage to the device. RESET according to the instructions. Failure to main board. Contact Technical Service (see Appendix). RESET according to the instructions. E004 If the problem occurs again, have the electrical mains checked by a technician. Disturbance on the electrical mains. If the electrical mains is equipped with a Continuity system, have the system checked by a technician. RESET according to the instructions. E007 One or more rear fans failed Check the operation of rear fans and contact Technical Service (see Appendix). RESET according to the instructions. If no integrated filters are equipped, empty the filling tank and refill it with distilled water of adequate quality (<15 µs/cm). The filling/discharge tank contains water of inadequate F008 quality. If an automatic filling system is present, empty the external container and fill it with water of adequate quality. If a Pure100/500 demineraliser or integrated filters are present, replace the filter elements. RESET according to the instructions. If no integrated filters are equipped, empty the filling tank IMMEDIATELY and refill it with distilled water of adequate quality (<15 µs/cm). If an automatic filling system is present, IMMEDIATELY empty the external container and fill it with water of adequate quality. If a Pure100/500 demineraliser or integrated The filling/discharge tank contains water of very poor filters are present, replace the filter elements IMMEDIATELY. F009 quality. In these conditions, the sterilizer allows starting a maximum of 5

consecutive cycles, after which it locks until the tank is filled with distilled water of adequate quality (<15 µs/cm) or integrated filters (if any) are replaced. This precaution is necessary to prevent damage

RESET according to the instructions. Empty both tanks IMMEDIATELY,

to the device.

Contact Technical Service

(see Appendix).

Door open (or not properly closed) at program start

The discharge tank contains water of very poor

Door position microswitch failure.

(START).

quality.

E010

E011

RESET according to the instructions.

Properly close the door and restart the program.

replace filter elements and fill the filling tank.

E012 The cycle limit after which integrated filters must be replaced has been reached. RESET according to the instructions. Empty that hanks IMMEDIATELY, replace filter elements and fill the filling tank.	CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
The filling tank contains water of very poor quality. and replace the demineralisation filter. In these conditions, the sterifizer allows sterring a maximum of S consecutive cycles, after which cycles are automatically symm. This procusion is necessary to prevent damage to the device.	E012		
E020 Door lock system gearmotor failure. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).	E013	The filling tank contains water of very poor quality.	and replace the demineralisation filter. In these conditions, the sterilizer allows starting a maximum of 5 consecutive cycles, after which cycles are automatically aborted until the correct water quality is detected (<15 µs/cm). This
Book lock system geatmost raisure. RESET according to the instructions. Contact Technical Service (see Appendix).	E020		Try restarting the program a second time.
Door lock system gearmotor failure. Contact Technical Service (see Appendix).		Door lock system gearmotor failure.	ii the presion poreiote, contact recrimical correct (coo the rippendix).
Bool lock microswitches failure Contact Technical Service (see Appendix).	E021		
Contact Technical Service (see Appendix). RESET according to the instructions. Top up with water up to the MAX level (or at least up to over the MIN level). RESET according to the instructions. Top up with water up to the MAX level (or at least up to over the MIN level). RESET according to the instructions and empty the tank. Contact Technical Service (see Appendix). RESET according to the instructions and empty the tank. Completely drain the drain tank. Completely drain the drain tank. Contact Technical Service (see Appendix). Reset according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Reset according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moisten		Door lock system gearmotor failure.	
Water level in the filling tank below minimum. Top up with water up to the MAX level (or at least up to over the MIN level). Contact Technical Service (see Appendix).	E022	Door lock microswitches failure	
Water level sensor failure. (see Appendix). RESET according to the instructions and empty the tank. Completely drain the drain tank. MAX water level sensor failure. Contact Technical Service (see Appendix). Problem in the hydraulic circuit. E042 Warning that the maximum water level in the tank has been reached (manual filling) Interrupt the filling operation to prevent water spillage. E050 Reminder to run the Vacuum Test cycle Run Vacuum Test as planned E051 Reminder to run the Helix Test cycle Run Vacuum Test cycle as planned E052 Reminder to run the combined Vacuum + Helix Test cycle as planned E063 The autoclave cannot connect to Lan network Check that the Lan network chosen for the connection is working properly. E060 The autoclave cannot connect to Wi-Fi network Make sure that configuration parameters of the Lan network are correct. Check that the Lan network chosen for the connection is working properly. E061 The autoclave cannot connect to Wi-Fi network Make sure that configuration parameters of the Wi-Fi network are correct. Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). E061 Air seepage through the gasket. E062 Reminder to run the program. E063 Contact Technical Service (see Appendix). E064 Air seepage through the gasket. E065 Contact Technical Service (see Appendix). E066 Reminder to run the combined Vacuum + Helix Test combined vycle as planned E066 Reminder to run the combined Vacuum + Helix Test cycle as planned E067 Reminder to run the combined Vacuum + Helix Test cycle as planned E068 Reminder to run the combined Vacuum + Helix Test cycle as planned E069 Reminder to run the combined Vacuum + Helix Test cycle as planned E069 Reminder to run the combined Vacuum + Helix Test cycle as planned E060 Reminder to run the vacuum test cycle E061 Reminder to run the vacuum test cycle E062 Reminder to run the vacuum test cycle E062 Reminder to run the vacuum tes	E030	Water level in the filling tank below minimum.	
E031 MAX water level in the train tank over the MAX level. Completely drain the drain tank. Contact Technical Service (see <u>Appendix</u>). E042 Warning that the maximum water level in the tank has been reached (manual filling) E050 Reminder to run the Vacuum Test cycle Run Vacuum Test as planned Reminder to run the Helix Test cycle Run Helix Test cycle as planned Reminder to run the Combined Vacuum + Helix Test cycle as planned Reminder to run the combined Vacuum + Helix Test cycle as planned The autoclave cannot connect to Lan network E060 The autoclave cannot connect to Lan network Make sure that configuration parameters of the Lan network are correct. Check that the run remanging the Wi-Fi network is and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. ResET according to the instructions.		MIN water level sensor failure.	
MAX water level sensor failure. (see Appendix).		Water level in the drain tank over the MAX level.	
E042 Warning that the maximum water level in the tank has been reached (manual filling) E050 Reminder to run the Vacuum Test cycle Run Vacuum Test as planned E051 Reminder to run the Helix Test cycle Run Vacuum Test as planned E052 Reminder to run the combined Vacuum + Helix Test cycle as planned E053 Reminder to run the combined Vacuum + Helix Test cycle as planned E054 Reminder to run the combined Vacuum + Helix Test cycle as planned E055 Reminder to run the combined Vacuum + Helix Test cycle as planned E066 The autoclave cannot connect to Lan network E067 The autoclave cannot connect to Lan network E068 The autoclave cannot connect to Wi-Fi network E069 The autoclave cannot connect to Wi-Fi network E069 The autoclave cannot connect to Wi-Fi network E060 Reminder to run the Vacuum + Helix Test cycle as planned Make sure that configuration parameters of the Lan network are correct. Check that the Instruction saverage of the Wi-Fi network is on and that the Wi-Fi network hosen for the connection is working properly. Contact Technical Service (see Appendix). E000 Reminder to run the Vacuum Test cycle E001 Air seepage through the gasket. E001 Reminder to run the Vacuum Test cycle E002 Reminder to run the Helix Test cycle as planned E003 Run Helix Test cycle as planned E004 Run Vacuum + Helix Test cycle as planned E005 Run Helix Test cycle as planned E005 Run Helix Test cycle as planned E006 Run	E031	MAX water level sensor failure.	
E042 has been reached (manual filling) Interrupt the filling operation to prevent water spillage.		Problem in the hydraulic circuit.	
Reminder to run the Helix Test cycle Run Helix Test cycle as planned	E042		Interrupt the filling operation to prevent water spillage.
Reminder to run the combined Vacuum + Helix Test cycle Run Vacuum + Helix Test combined cycle as planned Make sure that configuration parameters of the Lan network are correct. Check that the Lan network chosen for the connection is working properly. Contact Technical Service (see Appendix). The autoclave cannot connect to Wi-Fi network Make sure that configuration parameters of the Wi-Fi network are correct. Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program.	E050	Reminder to run the Vacuum Test cycle	Run Vacuum Test as planned
E060 The autoclave cannot connect to Lan network E061 The autoclave cannot connect to Wi-Fi network E061 Raw Surre that configuration parameters of the Wi-Fi network are correct. Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). E061 Raw Surre that configuration parameters of the Wi-Fi network are correct. Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). E061 Raw Surre that configuration parameters of the Lan network are correct. Check that the Lan network chosen for the connection is working properly. Contact Technical Service (see Appendix). E061 Raw Surre that configuration parameters of the Lan network chosen for the connection is working properly. Check that the Final Plan Plan Plan Plan Plan Plan Plan P	E051	Reminder to run the Helix Test cycle	Run Helix Test cycle as planned
E060 The autoclave cannot connect to Lan network Check that the Lan network chosen for the connection is working properly. Contact Technical Service (see Appendix). The autoclave cannot connect to Wi-Fi network	E052		Run Vacuum + Helix Test combined cycle as planned
E901 The autoclave cannot connect to Wi-Fi network Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit Contact Technical Service Contact Technical Service	E060	The autoclave cannot connect to Lan network	Check that the Lan network chosen for the connection is working properly.
E900 Air seepage through the gasket. Problem in the hydraulic circuit. Excessive humidity in the sterilization chamber. E901 Air seepage through the gasket. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Contact Technical Service (see Appendix). RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit. Contact Technical Service	E061	The autoclave cannot connect to Wi-Fi network	Check that the router managing the Wi-Fi network is on and that the Wi-Fi network chosen for the connection is working properly.
Excessive humidity in the sterilization chamber. Excessive humidity in the sterilization chamber. Excessive humidity in the sterilization chamber. RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit Contact Technical Service	E900	Air seepage through the gasket.	Thoroughly clean the gasket with a clean cotton cloth moistened with water.
Excessive numidity in the sterilization chamber. Thoroughly dry the inside of the chamber and restart the program. RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Problem in the hydraulic circuit Contact Technical Service		Problem in the hydraulic circuit.	
Air seepage through the gasket. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program. Contact Technical Service	E901	Excessive humidity in the sterilization chamber.	
Problem in the hydraulic circuit		Air seepage through the gasket.	Thoroughly clean the gasket with a clean cotton cloth moistened with water.
		Problem in the hydraulic circuit.	

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Excessive humidity in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program.
E902	Air seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program.
	Vacuum pump failure.	Contact Technical Service
	Problem in the hydraulic circuit.	(see Appendix).
E998	Service maintenance in progress.	Service maintenance in progress. If you were not informed, contact IMMEDIATELY the manager of the network to which the sterilizer is connected. Contact Technical Service (see <u>Appendix</u>).
E999	Manual interruption of the sterilization or test cycle.	RESET according to the instructions.

17.5.2. ALARMS (CATEGORY A)

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
A032	Connector of water level sensors in the filling tank not connected.	Contact Technical Service (see Appendix).
	Failure of water level sensor(s) in the filling tank.	
A033	Connector of water level sensors in the discharge tank not connected. Failure of water level sensor(s) in the discharge tank.	Contact Technical Service (see Appendix).
A034	Filter in the water filling tank removed during cycle	Refit the integrated filter inside the filling tank
A035	Filter in the water drain tank removed during cycle	Refit the integrated filter inside the discharge tank
	Lack of water in the external container (automatic filling)	RESET according to the instructions. Fill the container with a sufficient quantity of water (check the level at regular intervals).
A040	Automatic filling system not properly installed.	RESET according to the instructions. Check that the filling tube is properly connected. Remove any obstruction along the tube path.
	Automatic filling system failure.	Contact Technical Service (see Appendix).
A042	Possible problem to the Automatic filling system	Contact Technical Service (see Appendix).
A043	Possible problem in the Water recirculation system	Contact Technical Service (see Appendix).
A101	Chamber temperature sensor failure (PT1).	
A102	Steam generator temperature sensor failure (PT2).	
A103	Heating element temperature sensor failure (PT3).	
A105	Temperature sensor PT5 failed (conductivity measurement compensation)	
A111	Incorrect temperature sensor connection (sterilization chamber). Temperature sensor short-circuit (sterilization chamber).	
A112	Incorrect temperature sensor connection (steam generator).	Contact Technical Service (see Appendix).
	Temperature sensor short-circuit (steam generator).	
	Incorrect temperature sensor connection (heating	
A113	element). Temperature sensor short-circuit (heating element).	
A115	Temperature sensor PT5 short-circuited (conductivity measurement compensation).	
A116	ADC error.	
A120	Reference heating element acquisition chain fault.	
A121	Reference heating element acquisition chain fault.	
A122	Reference heating element acquisition chain fault.	Contact Technical Service (see Appendix).
A123	Solenoid valves control driver fault	
A124	Door motor control driver fault	
A125	Faulty current draw detected	Check mains voltage.
A126	Connection error with Wi-Fi module	Contact Technical Service (see Appendix).
A127	Communication error between graphical interface and process board through Can	
A128	Communication error between graphical interface and ethernet module	
A129	Connection error with NFC module NFC module failed	
A130	Connection error with RGB led bar RGB led bar failed	
A131	Solenoid valve 1 failed	Contact Technical Service (see Appendix).
A132	Solenoid valve 2 failed	
A133	Solenoid valve 3 failed	
A134	Solenoid valve 4 failed	
A135	Solenoid valve 5 failed	
A136	Solenoid valve 6 failed	
A137	Solenoid valve 7 failed	
A201	Steam generator safety thermostat triggered.	
	Steam generator or heating element malfunction.	
A202	Heating element safety thermostat triggered. Steam generator or heating element malfunction.	Contact Technical Service (see Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Water or condensate in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the sterilization chamber and restart the cycle. Do <u>not</u> insert material impregnated with water or in general with liquids into the chamber.
A250	Drain filter obstructed.	Clean the drain filter. (See Appendix Maintenance).
7,230	Air seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the cycle.
	Vacuum pump failure.	Contact Technical Service (see Appendix).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Water injection pump malfunction.	
A251	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Steam generator safety thermostat triggered. Steam generator malfunction.	
	Steam seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the cycle.
A252	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in <u>Appendix</u> Technical Characteristics).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	Contact Technical Service (see <u>Appendix</u>).
	Steam generator malfunction. Water or condensate in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the sterilization chamber and restart the program. Do <u>not</u> insert material impregnated with water or in general with liquids into the chamber.
A253	Air seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program.
	Vacuum pump failure.	
	Problem in the hydraulic circuit.	Contact Technical Service (see <u>Appendix</u>).
	Water injection pump malfunction.	
A254	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Steam generator safety thermostat triggered.	
	Steam generator malfunction. Steam seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program.
A255	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in <u>Appendix</u> Technical Characteristics).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	Contact Technical Service (see Appendix).
A256	Steam generator malfunction. Water or condensate in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the sterilization chamber and restart the program. Do <u>not</u> insert material impregnated with water or in general with liquids into the chamber.
	Air seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water. Restart the program.
	Vacuum pump failure.	Contact Technical Service (see Appendix).
	Problem in the hydraulic circuit.	Santaot roominour corrido (300 //pportaix).
A257	Water injection pump malfunction.	
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	Contact Technical Service (see <u>Appendix</u>).
		_
	Steam generator malfunction.	

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Steam seepage through the gasket.	RESET according to the instructions. Thoroughly clean the gasket with a clean cotton cloth moistened with water and restart the program.
A258	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in <u>Appendix</u> Technical Characteristics).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	Contact Technical Service (see <u>Appendix</u>).
	Steam generator malfunction.	
A260	Drain filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A200	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A261	Bacteriological filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A201	Problem in the hydraulic circuit.	Contact Technical Service (see <u>Appendix</u>).
A262	Bacteriological filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A202	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A050	Drain filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A353	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A050	Drain filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A356	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A260	Drain filter obstructed.	Clean the drain filter (see Maintenance Appendix).
A360	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A362	Drain filter obstructed.	Clean the drain filter (see Maintenance Appendix).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).

17.5.3. HAZARDS (CATEGORY H)

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
H150	Pressure sensor failure (MPX).	
H160	Pressure sensor (MPX) not properly connected to the connector.	
11100	Pressure sensor short-circuit (MPX).	
H400	Problem in the hydraulic circuit.	
H401	Problem in the hydraulic circuit.	
H402	Steam generator malfunction.	
11402	Problem in the hydraulic circuit.	
H403	Steam generator malfunction.	
11403	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
H404	Problem in the hydraulic circuit.	
11404	Steam generator malfunction.	
H405	Problem in the hydraulic circuit.	
11405	Steam generator malfunction.	
H406	Problem in the hydraulic circuit.	
П400	Steam generator malfunction.	
H410	Timer problem.	
H990	General operating problem.	
H991	General operating problem.	
H992	General operating problem.	
H993	General operating problem.	

17.5.4. SYSTEM ERRORS (CATEGORY S)

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
S001	Error of Flash memory 1 on process board Flash memory 1 on process board failed	Contact Technical Service (see Appendix).
S002	Error of Flash memory 2 on process board Flash memory 2 on process board failed	Contact Technical Service (see Appendix).
S003	Error of flash memory 1 on graphical interface board Flash memory 1 on graphical interface board failed	Contact Technical Service (see Appendix).
S004	Error of flash memory 2 on graphical interface board Flash memory 1 on graphical interface board failed	Contact Technical Service (see Appendix).
S005	USB key not correctly formatted Damaged USB key	Check USB key correct formatting (FAT32). As an alternative, use another correctly formatted USB key. If the problem persists, contact Technical Service (see Appendix).
S006	USB key not correctly formatted Damaged USB key	Check USB key correct formatting (FAT32). As an alternative, use another correctly formatted USB key. If the problem persists, contact Technical Service (see Appendix).
S007	USB key full	Download data from USB key or use another USB key. If the problem persists, contact Technical Service (see Appendix).
S009	Printer off. Data cable not correctly connected to serial ports RS-232.	Make sure that printer is on. Check printer cable correct connection. If the problem persists, contact Technical Service (see Appendix).
S010	No paper inside printer. Paper setting configuration not correctly done.	Make sure that paper is correctly loaded. Check printer cable correct connection. Make sure that paper settings are correct. If the problem persists, contact Technical Service (see Appendix).
S011	Printer lid open	Make sure that printer lid is correctly closed. Check printer cable correct connection. If the problem persists, contact Technical Service (see Appendix).
S012	Printer not ready for use	Make sure that paper is correctly loaded. Check printer cable correct connection. Make sure that paper settings are correct. If the problem persists, contact Technical Service (see Appendix).
S020	Cycle back-up not done after 250 cycles	Perform cycle back-up. See paragraph Sterilization cycle back-up. If the problem persists, contact Technical Service (see Appendix).
S021	Cycle storage limit exceeded after 500 cycles	Perform cycle back-up. See paragraph Sterilization cycle back-up. If the problem persists, contact Technical Service (see Appendix).
S030	Malfunction of the control software	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
S031	Malfunction of control board or software	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
S032	Malfunction of the control software	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
S034	Malfunction of the control software	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
S035	Control software malfunction in solenoid valve management	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
S040	Malfunction of the control software	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
S041	Malfunction of control board or control software	Contact Technical Service (see Appendix).
S042	Malfunction of control board or software	Contact Technical Service (see Appendix).
S099	Malfunction of control board or control software	Try restarting the program a second time. Try replacing the USB key. If the problem persists, contact Technical Service (see the Appendix).
S100	Malfunction of control board or control software	Contact Technical Service (see Appendix).

18. USER PIN RESET



If the user enters the pin incorrectly for 3 times, it is necessary to enter the following unlock pin for four consecutive times when you will be prompted to enter pin again:

9999

19. APPENDIX - ACCESSORIES

Only use spare parts and accessories that meet the manufacturer's specifications.

DEMINERALIZER PURE 100

DEMINERALIZER PURE 500 TWIN PURE 500

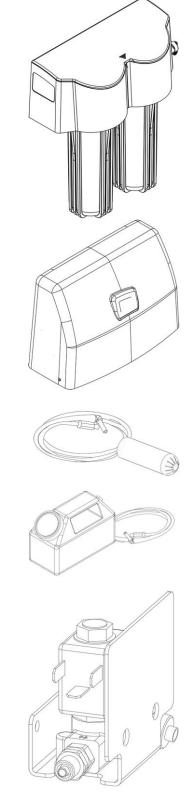
AUTOMATIC FILLING

FRONT FILLING

H₂O AUXILIARY SOLENOID VALVE

Additional SV kit including:

- 2-way water solenoid valve, NC 24 V DC 1
- Steel support and fastening screws
- Connection cable with plug 3
- Silicone hose with connector
- 5 Control valve
- 1-way valve

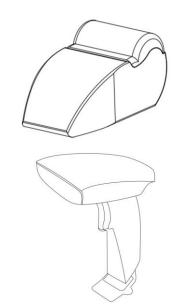




For the management of automatic filling accessories, refer to the manual of the relevant accessory.

EXTERNAL PRINTER

BARCODE READER



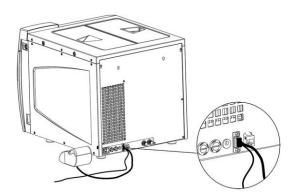
DATA STER SOFTWARE

MY TRACE SOFTWARE

20. PRINTER CONNECTION

Connect the printer to the RS232 serial port located on the rear of the autoclave (see figure).

Load the desired type of paper and turn on the printer. Set the type of paper loaded (see the paragraph PRINT MANAGEMENT).





Refer to the printer manual for printer starting and paper loading.

21. APPENDIX - SPARE PARTS AND ACCESSORIES

Only use spare parts and accessories that meet the manufacturer's specifications.

Description	Code	
bacteriological filter	97290160	
door gasket (17/22 I)	97400145	
door gasket (28 I only)	97467176	
demineralised water tank/chamber filter	97290210	
integrated demineralisation filter	97290259	

22. APPENDIX - TECHNICAL SERVICE

FOR ANY REQUEST FOR TECHNICAL INTERVENTION ON THE PRODUCT,
BOTH UNDER WARRANTY AND OUT OF WARRANTY, DIRECTLY CONTACT
THE DEALER OR RESELLER THAT SUPPLIED IT.

We will gladly provide any information you may need on the product as well as give you suggestions and advice on the water steam sterilization procedures. In this regard, please refer to the following address:

Cefla S.c.

Plant Via Bicocca, 14/C 40026 - Imola (BO) IT Tel. +39 0542 653441 Fax. +39 0542 653555

Headquarters Via Selice Provinciale 23/A – 40026 Imola (BO) IT

23. APPENDIX - WARNINGS AND LOCAL REGULATIONS

Please consult the Web site of the manufacturer to find a list of authorised representatives.



Before carrying out any technical service operations, consult the service manual containing the above instructions.

