97055023
Rev. 03
2021-03

Supreme-17 / Supreme-22 / Supreme-28

CEFLA S.C. VIA SELICE PROVINCIALE 23/A - 40026 IMOLA (BO) ITALY
PLANT: VIA BICOCCA 14/C - 40026 IMOLA (BO) - ITALY

EN

Contents

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.

CE 0051
Device in compliance with essential requirements of Directive 93/42/EU and subsequent modifications.
Notified body: IMQ spa

Refer to the user manual.

CE 0474
Device compliant with the requirements set out in the Directive 2014/68/EU (PED) - category I for 17 l sterilizers; category II for 22 and 28 l sterilizers.
Notified body: Rina Services S.p.A.

UA.TR.101
Ukrainian national symbol of conformity.

I / O
ON/OFF switch.

2xT15A 250V
Fuses 2xT15A 250V.

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.
2014/68/EU, (**PED**).

The product complies with Standard **EN 13060:2014 + A1:2018.**

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.

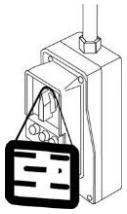
Please observe the following precautions in order to avoid injury or property damage:

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

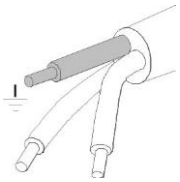
The use of water of inadequate quality can severely damage the device.
See technical characteristics appendix in this regard.

- **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**.



Failure to comply with the above exempts the manufacturer from all liability.

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.

2. PACKAGE CONTENT

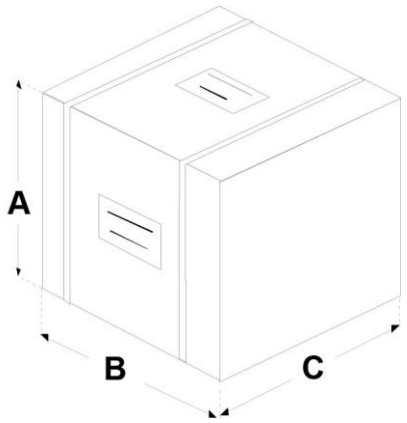
Check the integrity of the product package upon receipt.

2.1. DIMENSIONS AND WEIGHT

Once the package is opened, check that:

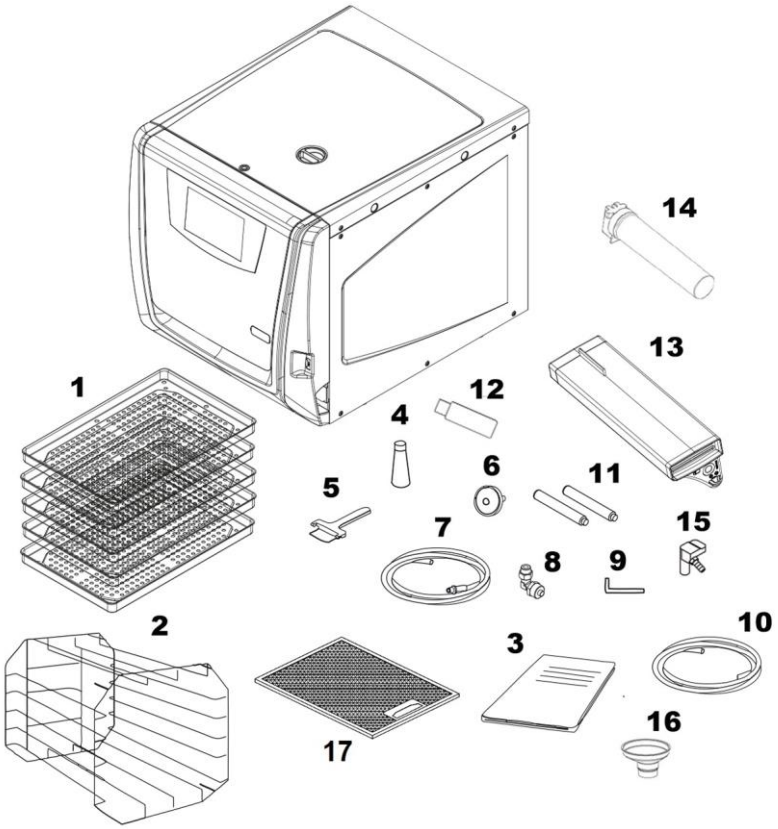
- The supply matches the specifications of the order (see the delivery note);
- There is no visible damage to the product.

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:

1 Instrument trays:
- 5 pcs for 17 and 22
- 6 pcs for 28;

2 Tray holder support;

3 Operator's documentation and safety valve's EC Declaration of Conformity;

4 Lubricant for door locking mechanism;

5 Tray extractor;

6 Additional bacteriological filter.

7 Rubber hose with quick coupling for manual water drainage;

8 Angled union + straight union;

9 Allen wrench (for door manual unlocking);

10 Plastic tube for direct water drainage, with fastening clamp;

11 Rear spacers;

12 USB key containing:
User manual and DataSter Software;

13 Filling tank demineralisation filter;

14 Discharge tank recirculation filter;

15 Discharge tank recirculation filter union
(to be used when recirculation filter is not present);

16 Funnel for water loading.

17 Dust filter.

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 distilled water and used water 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 0.5 and 1.1 bar (50 and 110 kPa)

3. 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.

A lighting system is activated when the door is opened for the working area in front of the sterilization chamber, allowing performing daily operations with increasing ease and comfort. There is also a remote light signalling device integrated into the front panel of the device.

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

Device	WATER STEAM STERILIZER		
	17	22	28
Class (according to Directive 93/42/EEC and subsequent amendments)	IIb		
Manufacturer	CEFLA s.c. Headquarters Via Selice Provinciale 23/A – 40026 Imola (BO) IT		
Input voltage	220 V - 240 V~ 50 Hz 220 V - 240 V~ 60 Hz 120V~ 60 Hz		
Network fuses (6.3 x 32 mm)	2x T15A 250V		
Electronic board fuses (5 x 20 mm)	F1: T3.15A 250V (transformer primary 220/240 V~ 50 Hz 220/240 V~ 60 Hz) F2: T3.15A 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		
Operational environment	Indoor use HUMID LOCATION (EN 61010 extended environmental conditions)		
A-weighted sound power level (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.)		
External dimensions (HxWxD) (rear connections excluded)	500 x 480 x 600 mm		
Net weight: unladen unladen, with tray holder support and trays unladen, with tray holder support, trays and water at MAX level	approx. 51 kg approx. 54 kg approx. 60 kg	approx. 52 kg approx. 55 kg approx. 61 kg	approx. 53 kg approx. 56kg approx. 62 kg
Sterilization chamber dimensions (D x D)	250 x 350 mm	250 x 450 mm	280 x 450 mm
Sterilization chamber total volume	approx. 17 l (0.017 cu. m)	approx. 22 l (0.022 cu. m)	approx. 28 l (0.028 cu. m)
Sterilization chamber usable volume (with tray holder support inserted)	approx. 10 l (0.010 cu. m)	approx. 13 l (0.013 cu. m)	approx. 19 l (0.019 cu. m)
Sterilization chamber usable dimensions	17 l (1.38x1.55x2.97) dm / 6.4 cu. dm	22 l (1.38x1.55x3.97) dm / 8.5 cu. dm	28 l (1.72x1.66x3.96) dm / 11.3 cu. dm
Distilled water tank capacity (filling)	approx. 5.5 l (water at MAX level) approx. 1 l (water at MIN level)		
Sterilization programs	5 standard programs + 1 program defined by the user		
Test programs	Helix/BD TEST Vacuum Test Vacuum Test+Helix/BD Test		
Pre-heating time (from cold)	approx. 10 min		

Device	WATER STEAM STERILIZER		
	17	22	28
USB connection	Stick 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:	Compliant with Standard EN 60950. (The safety of the sterilizer may be compromised in case of uncertified printer power supply unit)		
120V 60Hz Main power cord:	Plug NEMA 5-15 125V-15A Cable SJT 14 AWG/3C STYLE 1015 60°C Connector C19 acc. to IEC 60320		
220-240V 50Hz Main power cord:	Plug CEE 7/VII IEC 250V-16A 50 Hz Cable 3x1.5 mm² -25 to 70 °C Connector C19 acc. to IEC 60320 UL 498, CSA C22.2		
220V 60Hz Main power cord:	Plug NEMA 6-15P 250V-15A SJT 14 AWG/3C 300V 60°C Connector C19 acc. to IEC 60320		
Ethernet connection	RJ45 (max. cable length 29 m)		
Wi-Fi	802.11 b/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 90° C		
Total heat in Joule sent by the sterilizer to the surrounding air in 1 hour of continue operation	17 l = 3.6 kJ	22 l = 4 kJ	28 l = 5.4 kJ
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	500 kPa (abs)	500 kPa (abs)	500 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.

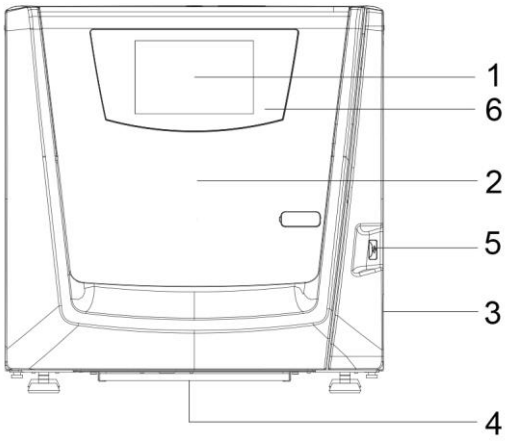
DESCRIPTION	VALUES IN THE WATER SUPPLY	VALUES INSIDE RESIDUAL
DRY CONDENSATE	< 10 mg/l	< 1 mg/l
SILICON OXIDE SiO_2	< 1 mg/l	< 0.1 mg/l
IRON	< 0.2 mg/l l	< 0.1 mg/
CADMIUM	< 0.005 mg/l	< 0.005 mg/l
LEAD	< 0.05 mg/l	< 0.05 mg/l
HEAVY METAL RESIDUES (iron, cadmium and lead excluded)	< 0.1 mg/l	< 0.1 mg/l
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.

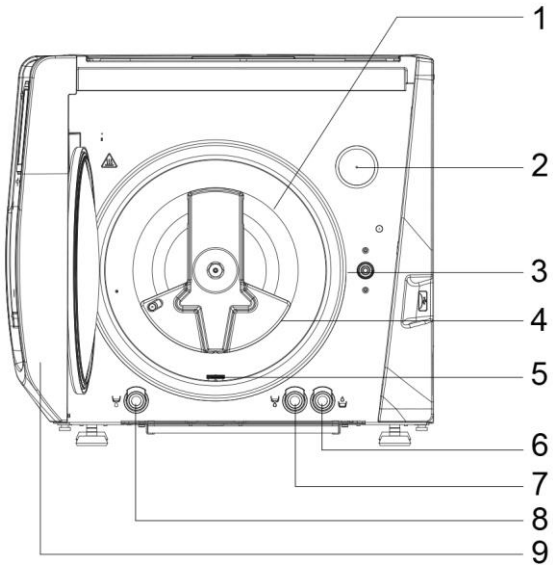
The use of water for steam generation with contaminant levels exceeding those indicated in the above table can greatly shorten the sterilizer lifetime.
This could also result in an increase of oxidation in the most sensitive materials as well as in an increase of limescale residues on generator, boiler, internal supports, trays and instruments.

3.5. FRONT

1 LCD touch screen control panel
2 Door
3 Power switch
4 Dust filter
5 USB port
6 NFC reader

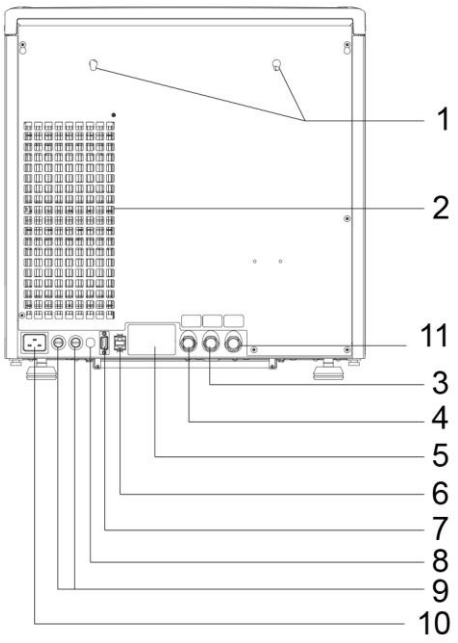


1 Sterilization chamber
2 Bacteriological filter
3 Door locking system
4 Steam diffuser
5 Water drainage filter
6 Water top-up quick connector
7 Water drainage quick connector
8 Waste water drainage quick connector
9 Door

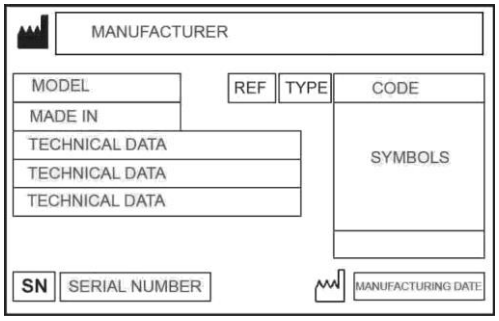


3.6. REAR

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, EV AUX kit and external pump 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, EV AUX kit and external pump kit)
9 Network fuses
10 Power cable connection
11 Connection for “overflow” drain



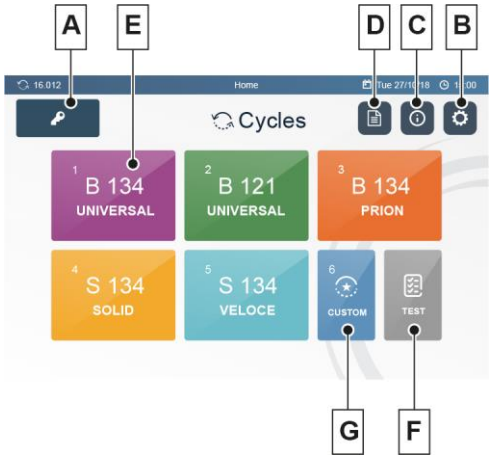
(*)



3.7. DISPLAY 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.

A Door opening
B Settings
C Sterilizer information
D Performed cycle list
E Sterilization cycles
F Test cycles
G Custom cycle (customisable)



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

3.8. EXAMPLE OF WORKING CYCLE

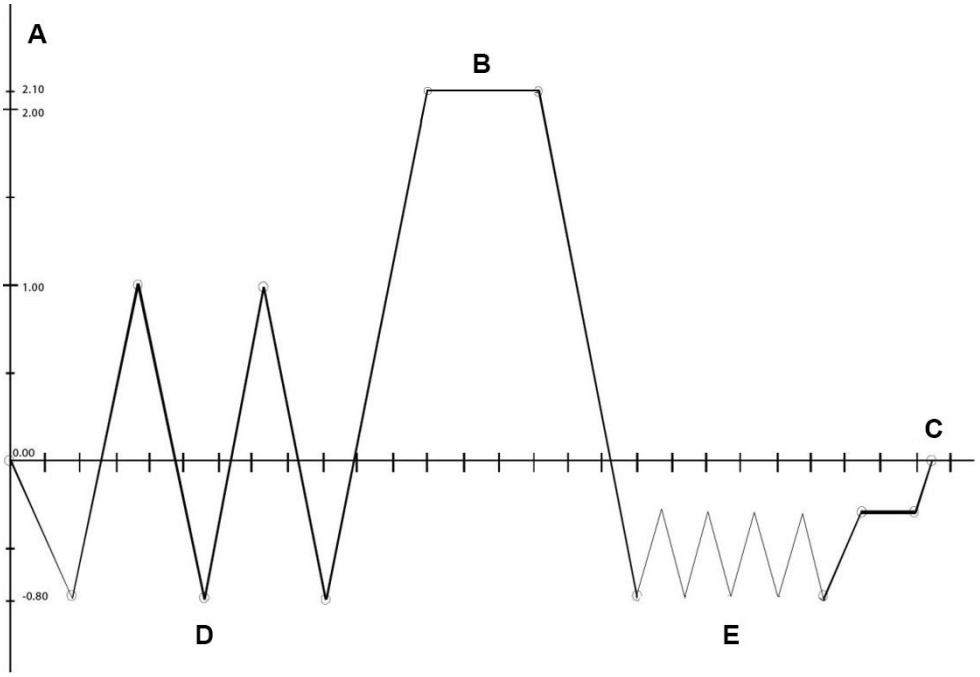
The sterilization program can be effectively described as a succession of phases, each one with a very precise objective.

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):

1 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;
3 Raising the pressure, with the consequent increase in the temperature of the steam, until reaching the conditions required for sterilization (in the example, 134°C);
4 Stabilizing the pressure and temperature;
5 Sterilizing for the required time (in the example, 4 minutes);
6 Depressurizing the sterilization chamber;
7 Vacuum-drying phase;
8 Ventilating the load with sterile air;
9 Bringing the pressure of the sterilization chamber back to the atmospheric level.

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.



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

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

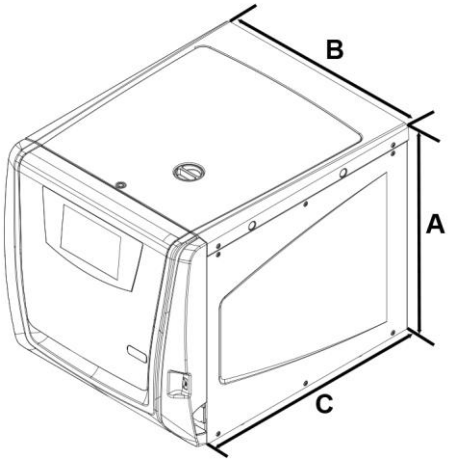
4. SETTING UP THE DEVICE

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.

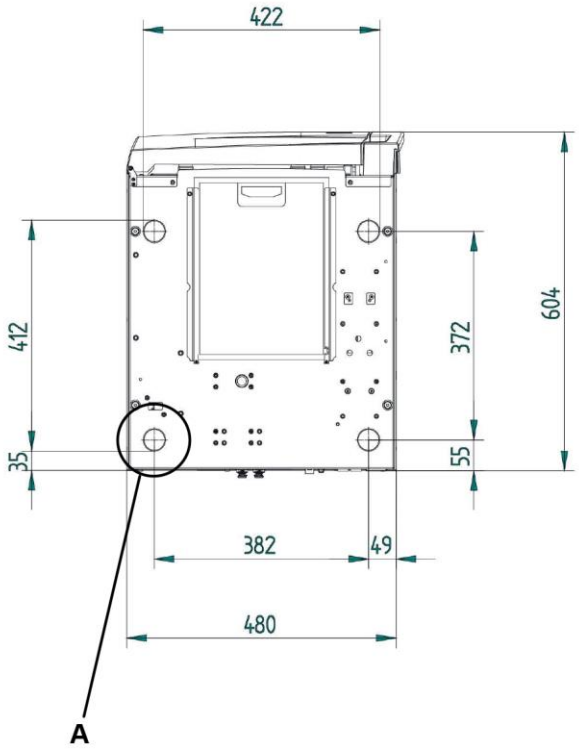
Technical service department (see appendix) is available for any doubt or further information. The sterilizer is placed on the marked only after having passed all the checks required. It does not require any additional calibration for commissioning.

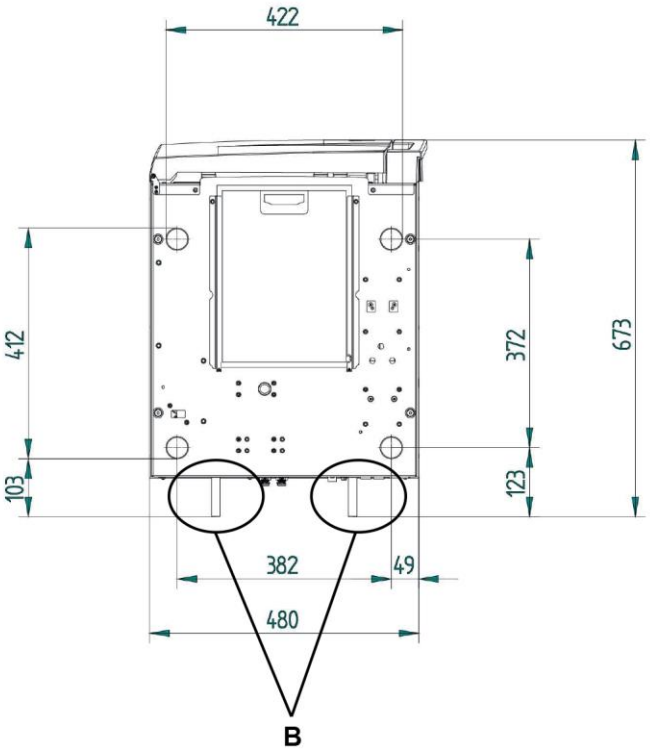
Dimensions and weight	**17 l**	**22 l**	**28 l**
A Height (total)	500 mm		
B Width (total)	480 mm		
C Depth (excluding rear unions). **Note: in any case, the sterilizer can be positioned on a plane of only 550**	600 mm		
Total weight	54 kg	55 kg	56 kg



4.1. OVERALL DIMENSIONS

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





A Feet
B Rear spacers

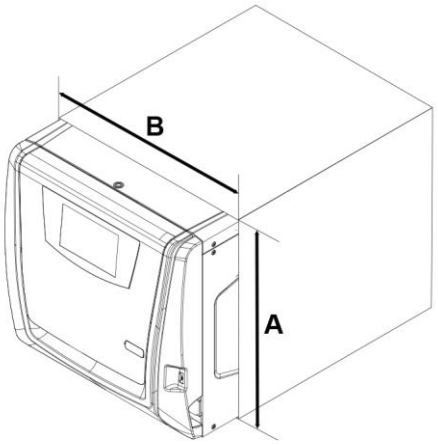
4.2. COMPARTMENT DIMENSIONS FOR BUILT-IN INSTALLATION

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
Height	520 mm WITH FRONT FILLING OR AUTOMATIC FILLING KIT
Width	550 mm
Depth	600 mm



In case of water manual filling from the top, the installation compartment must be equipped with an extractable table with suitable load bearing capacity (approx. 90Kg).

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

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 **16 A**
- 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.

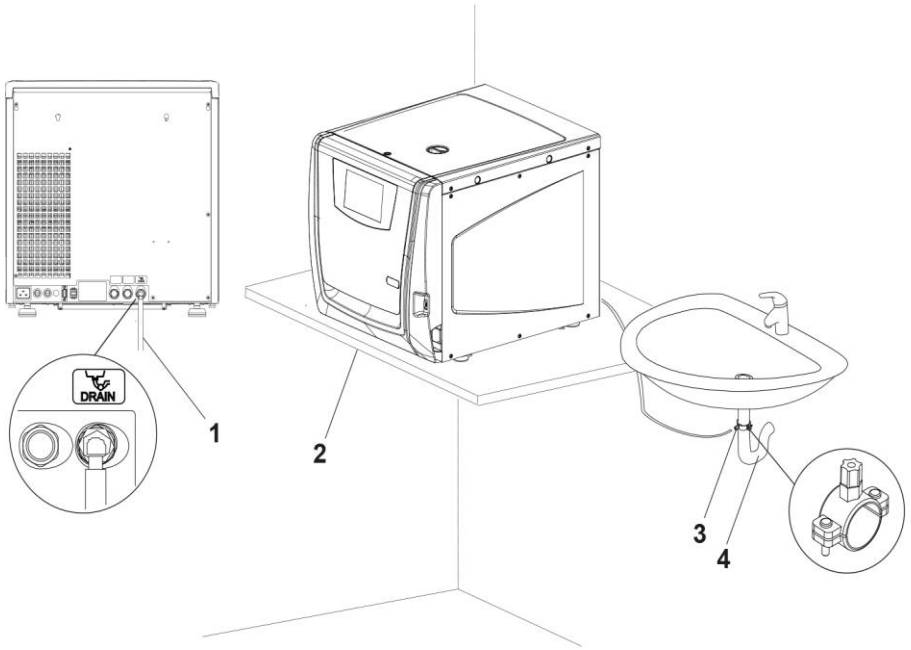
4.6. DIRECT CONNECTION TO A CENTRALIZED DRAINING POINT

Use direct discharge connection ONLY when using a device without discharge tank recirculation filter and filling tank demineralisation filter.

- 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;	**3**	Clamp;
2	Resting surface;	**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.

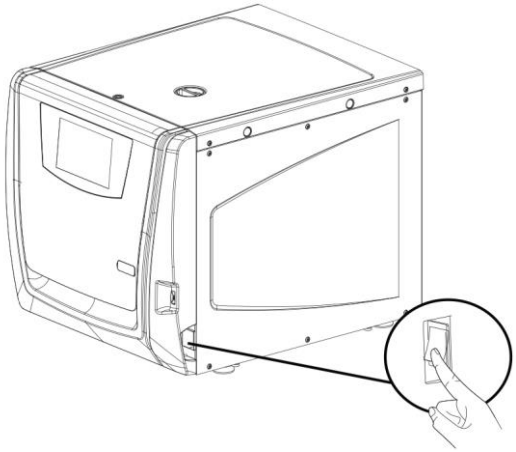
If an automatic filling system (external pump or solenoid valve, Pure 100, Pure 500) is connected, it is essential to use an overflow discharge connection.
In case of fault or failure, this system allows any excess water produced by the automatic filling system to flow into the centralized draining point,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

5. FIRST START-UP

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

5.1. STARTING

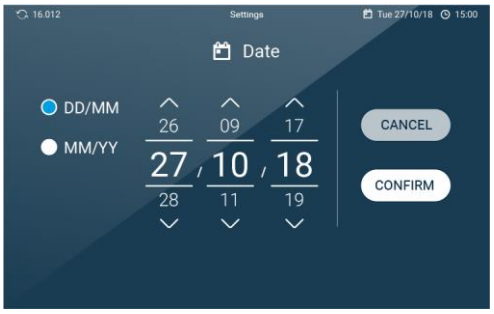
Once the sterilizer has correctly been installed, turn it on with the main switch on the right-hand side of the machine.

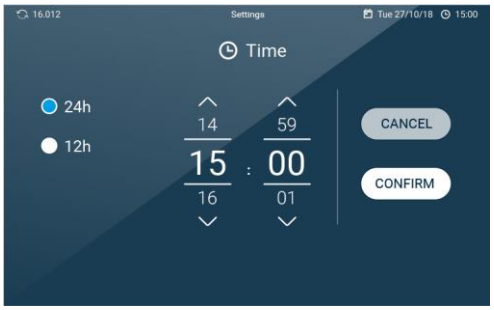


Do not turn on the sterilizer if USB key is inserted.

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.



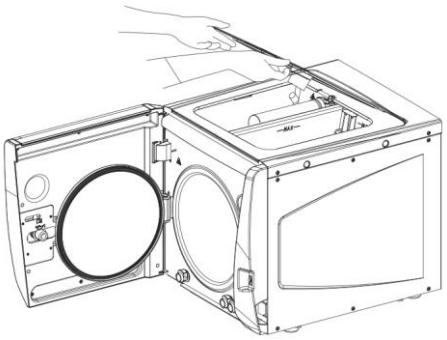


5.2. MAIN MENU

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

5.3. UPPER DOOR OPENING

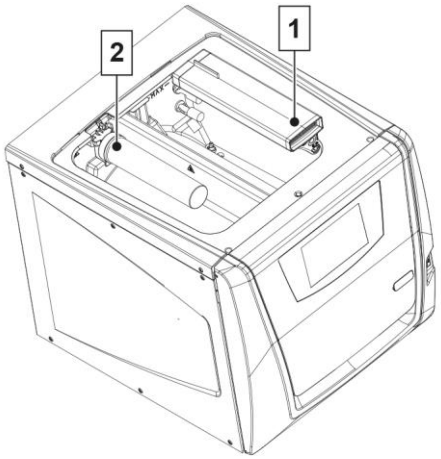
Remove the upper panel by lifting it and sliding it forward.



5.4. DISCHARGE WATER RECOVERY AND DEMINERALIZATION SYSTEM

The demineralisation and discharge water recovery system is an innovative instrument designed to treat mains water and recover the condensate drained from the sterilizer.
Filters are made of bacteriostatic material to prevent the proliferation of bacteria on their surfaces.
The system consists of two filters housed, respectively, inside the filling tank and inside the discharge tank:

1 FILLING TANK DEMINERALISATION FILTER
2 DISCHARGE TANK RECIRCULATION FILTER



The demineralisation filter is used to demineralise mains water, thanks to the use of ion-exchange resins. The system supplied distinguishes itself from other similar products on the market thanks to the fact that its installation requires no additional space, since it is housed inside the sterilizer, allowing the user to save space inside the practice.
The recirculation filter is used to recondition sterilizer's discharge water, making it chemically and microbiologically pure, so that it can be reused to generate steam.
This new feature allows saving up to 90% water as compared with a standard class B autoclave.
All this with the objective of offering a product suitable for the daily needs in the medical area.

The sterilizer will start using reconditioned water after the minimum level inside the discharge tank has been reached.

It is not necessary to empty the discharge water tank, except after a long machine downtime and when the recirculation filter is replaced.

Do not remove the demineralisation filter and/or recirculation filter during the sterilizer operation.

The sterilizer has been designed to operate with both filters fitted into the machine. The recirculation water cannot be used if the demineralisation filter and/or the recirculation filter are removed.

For a good operation of the demineralisation and recirculation filters, it is essential to install them correctly; in this way, it will be possible to prevent malfunctions or damage to the device. Please, strictly follow the warnings below.

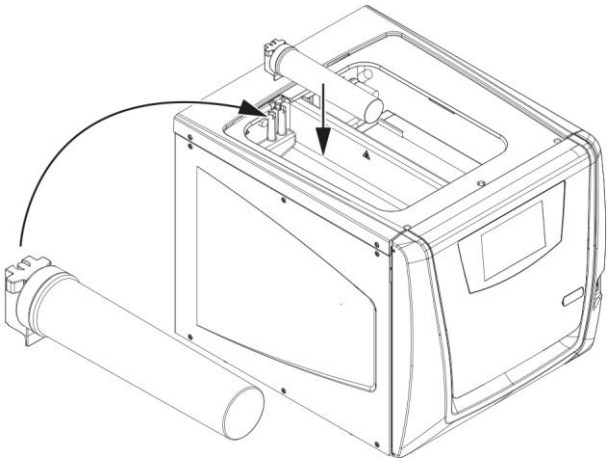
Install the demineralisation and recirculation filter only on dedicated sterilizers.
The technical assistance service is at your disposal for further information or any doubt you may have

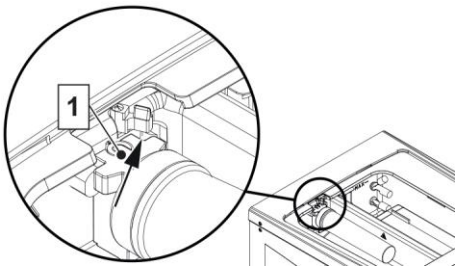
5.4.1. INSTALLING THE RECIRCULATION FILTER INSIDE THE TANK

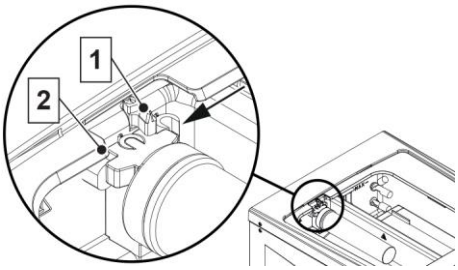
Take the recirculation filter from the package, then proceed as follows:

1 Turn the sterilizer off;
2 Move the filter locking lever (1) backward;
3 Fit the recirculation filter (2) onto the hole located in the upper part of the discharge tank;
4 Move the locking lever (1) forward until it reaches the upper part of the filter;
5 The sterilizer is ready to reuse the discharge water.

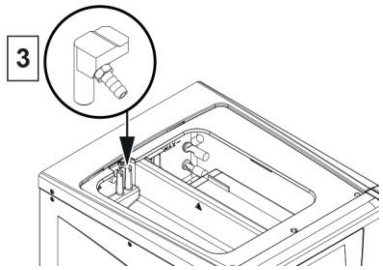
Both tanks must be emptied from water before leaving the sterilizer off for a long period of time.
Failure to empty them will cause the degradation of the water quality and the proliferation of bacteria.
Due to water stagnation in the demineralisation and recirculation filter ducts and in the sterilizer's tanks.

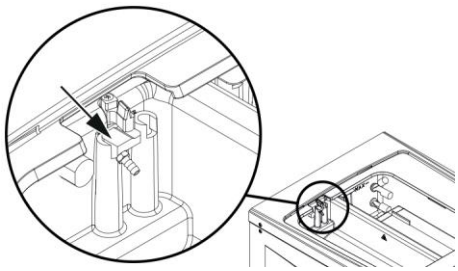






If the recirculation filter is removed and no longer used, fit in its place the union (3) supplied with the sterilizer, following the same procedure used to fit the recirculation filter inside the tank.



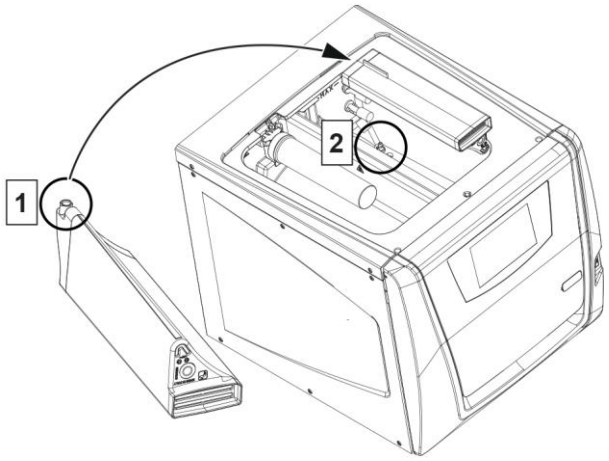


5.4.2. INSTALLING THE DEMINERALIZATION FILTER INSIDE THE TANK

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.

Take the demineralizer from the package, then proceed as follows:

1 Empty the water tank, if equipped, through the special pipe;
2 Fit hole (1) onto the metal union (2) at the bottom of the sterilizer's tank and exert a slight pressure;



3 Push the front side of the demineralizer downward until a "clack" is heard, indicating 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.

5.4.3. SAFETY PRECAUTIONS

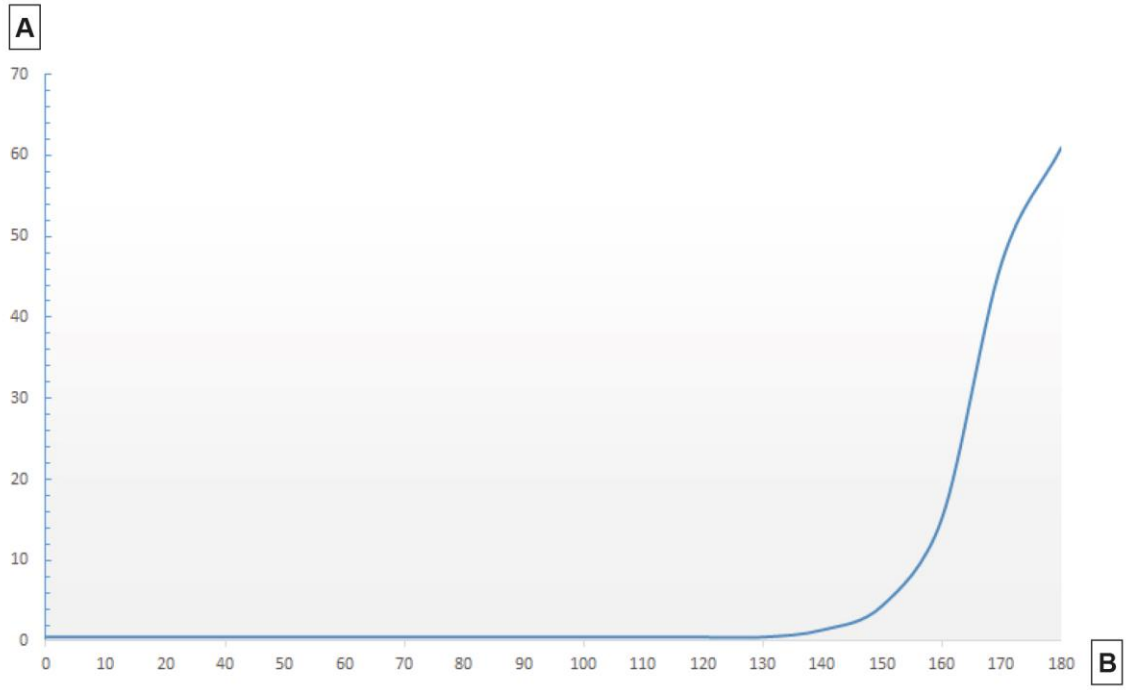
Below are the indications to be followed in case of contact with the ion-exchange resins contained in the cartridges of the demineralizer.

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.
	Remove all particles and rinse affected areas with water.
ACCIDENTAL DISPERSION MEASURES	
Precautions for people.	Keep people away.
	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	

5.4.4. TECHNICAL SPECIFICATIONS

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. 170 litres

Demineralisation filter performance



A H2O conductivity at filter outlet

B Litres of treated water

5.5. 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.

Upon the first use of the sterilizer and whenever a water shortage is signalled, fill or top up the water tank.

5.6. FILLING DISTILLED WATER

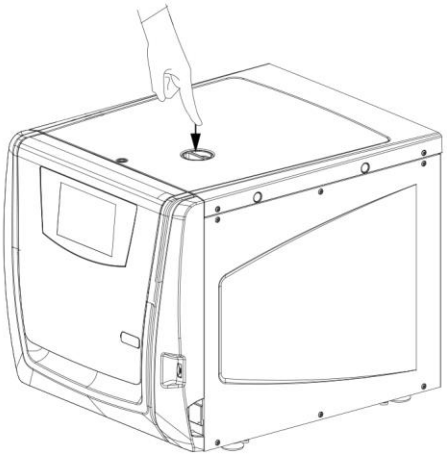
The sterilizer is equipped with a demineralisation filter fitted inside the filling tank, which allows supplying the device with standard water from the mains. The quality of the water treated by the integrated filter is checked automatically through a conductivity sensor.
If the demineralisation filter 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 and lead excluded)	< 0.1 mg/l	< 0.1 mg/l
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

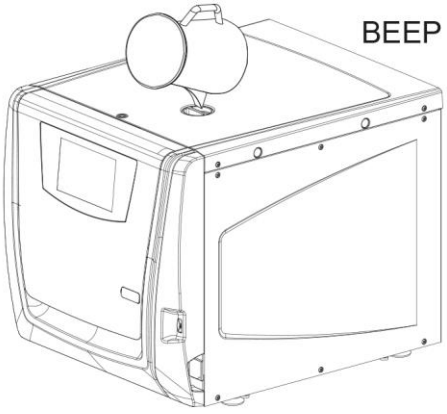
If the autoclave is to be used with distilled water, remove the integrated demineralisation filter from the filling tank.

5.6.1. MANUAL FILLING

Manual filling is performed through the filling hole fitted on the upper door, using mains water. Remove the plug from the hole and insert the supplied funnel.
Pour water through the funnel, making sure not to exceed the maximum level inside the tank (max).



When the MAX level is reached, the sterilizer will emit a sound.



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.

5.6.2. AUTOMATIC FILLING

Refer to the chapter WATER FILLING.

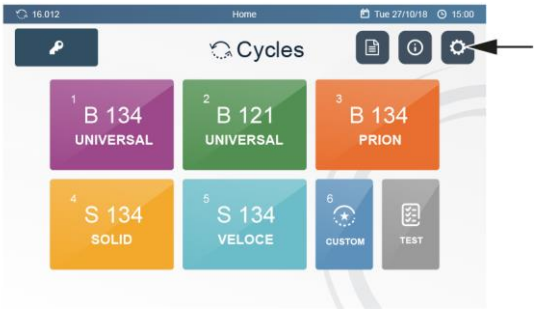
6. CONFIGURATION

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.



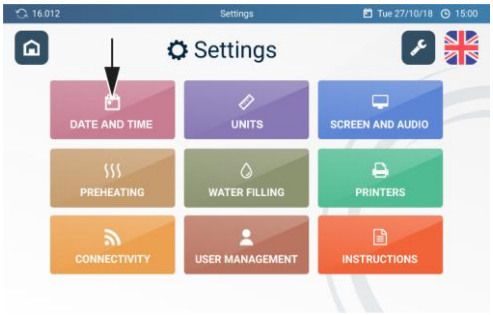
6.1.1. LANGUAGE

Select LANGUAGE.

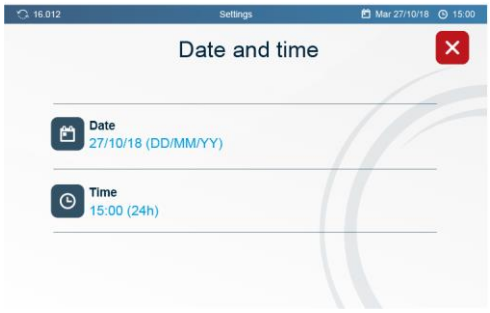
Select the desired language by scrolling the list and confirm by pressing the CONFIRM icon.

6.1.2. DATE AND TIME

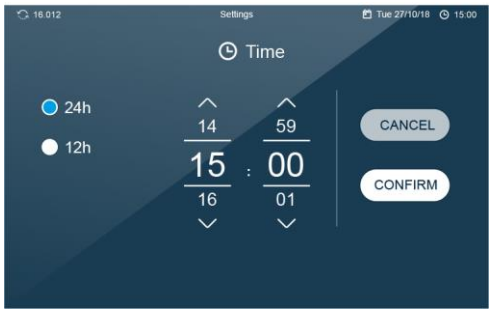
Select DATE AND TIME option.



Select the field to edit.



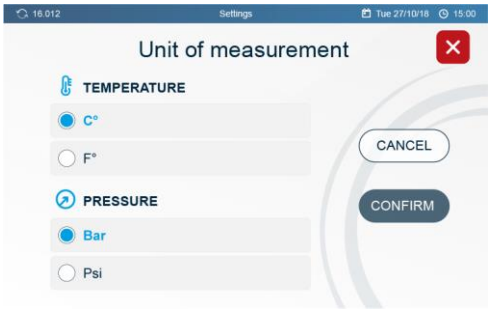
Set DATE and TIME and confirm by pressing the CONFIRM icon.



6.1.3. UNIT OF MEASUREMENT

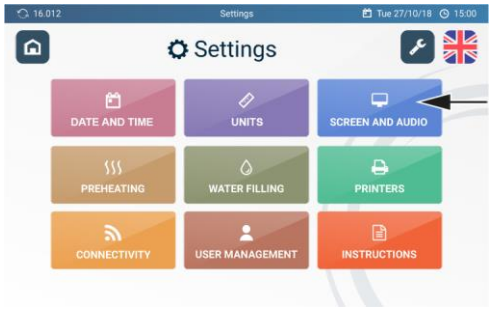
Select UNIT OF MEASUREMENT to set the reading of temperature and pressure values.

Confirm by pressing the CONFIRM icon

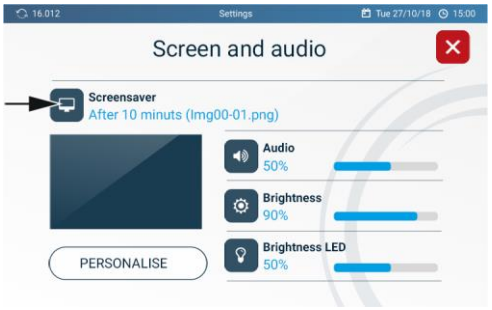


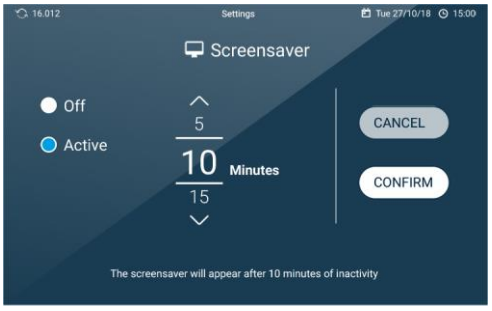
6.1.4. SCREEN AND AUDIO

Select SCREEN AND AUDIO to set the parameters of screensaver, audio volume and brightness of screen and LED bars.

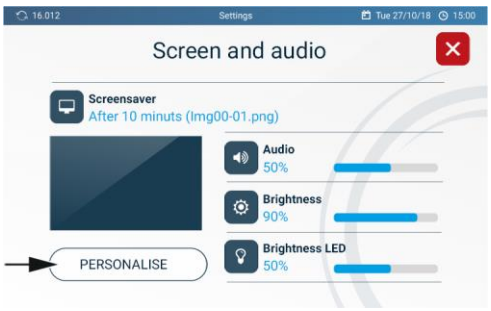


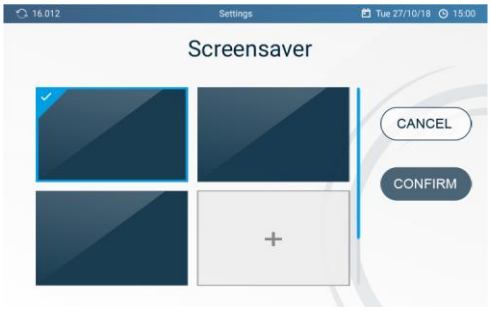
Use the Screensaver control to set the activation time.

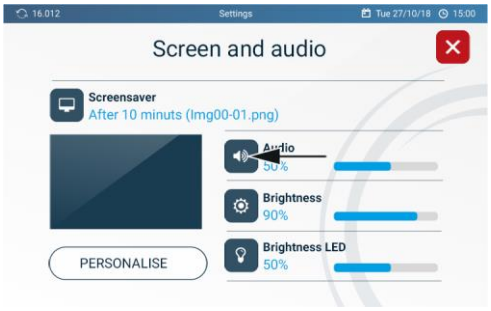




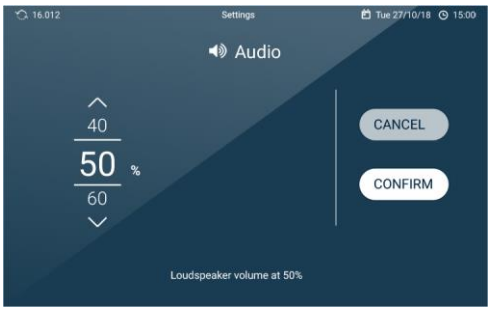
Use the Personalise control to select one of the screensavers provided or to load an image from an USB memory support to be connected to the front USB port.

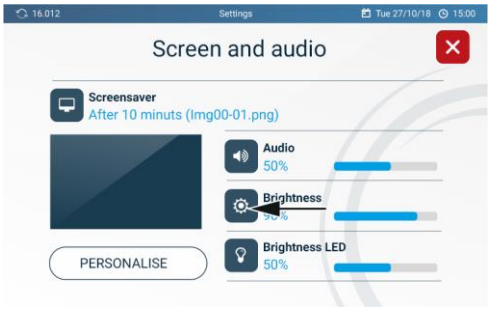




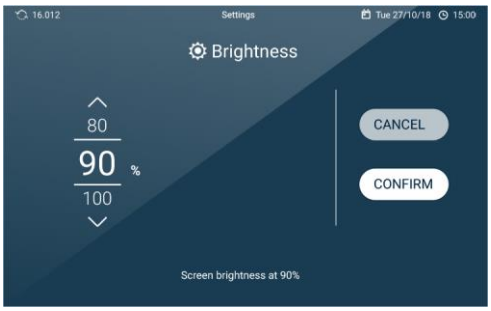


Use the AUDIO control to set the volume of notifications.



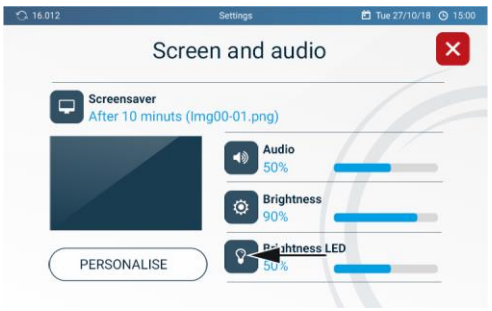


Use the BRIGHTNESS control to set the brightness intensity of the screen.



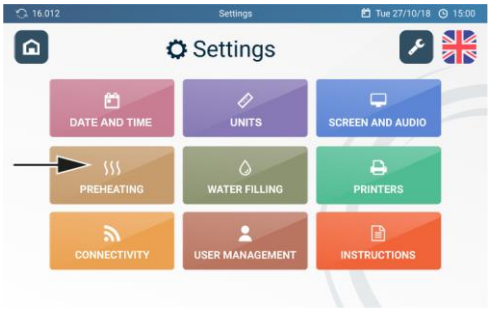
The LED Brightness control affects the internal LED bar, activated when the door is opened.

When the LED brightness value is set to 00, both bars remain off.

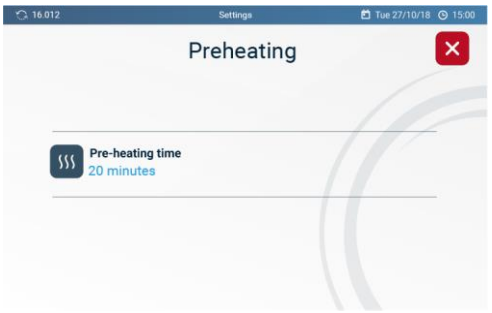


6.1.5. PREHEATING

Select PREHEATING to activate the PREHEATING of the sterilization chamber.



Select ACTIVE to activate the PREHEATING, OFF to disable it.



It is possible to set the maximum operating time after which the warming up is disabled.

6.1.6. WATER FILLING

Select WATER FILLING to choose the type of water filling.

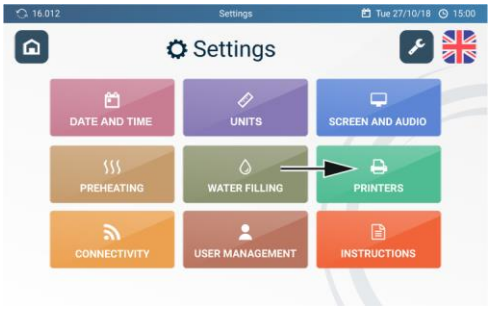
Available options include:

When the automatic filling system is connected, the sterilizer asks to Identify the type of device actually connected by pressing the corresponding key.
If connecting the filling system when the sterilizer is off, access the menu via the configuration program and manually select the correct option.

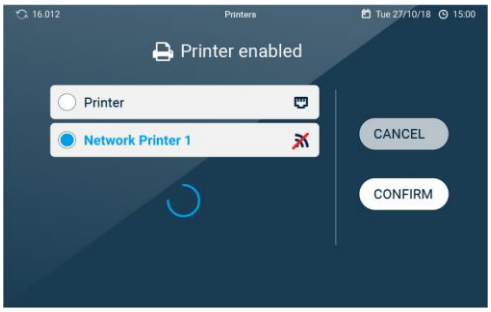
This menu can also be used to temporarily deactivate the automatic filling system (filters exhausted, fault, etc.) and go to manual tank filling, keeping the automatic filling system connected.

6.1.7. PRINTERS

Select PRINTERS to choose which support to use for printing.



Select PRINTER ENABLED to view the list of printers that can be controlled from the device, which are connected via RS232 port or Wi-Fi.

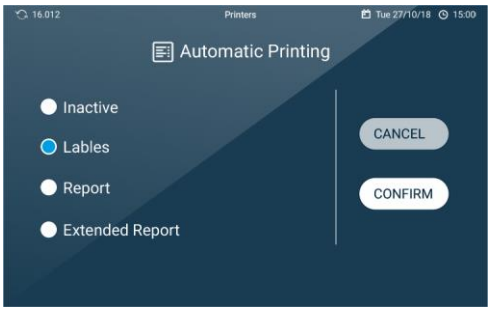


Select AUTOMATIC PRINTING to access the print mode selection menu, activated when the door is opened at the end of the cycle.

- INACTIVE - no printing at the end of the cycle.
- LABELS - label printing activated, including barecodes, cycle data and device data.
- REPORT - printing of a compact version of the cycle summary report activated.
- EXTENDED REPORT - printing of an extended version of the cycle summary report dedicated to the Technical Service activated.

Once the print mode has been selected, check which type of support (thermal paper roll, label roll) is loaded onto the printer.

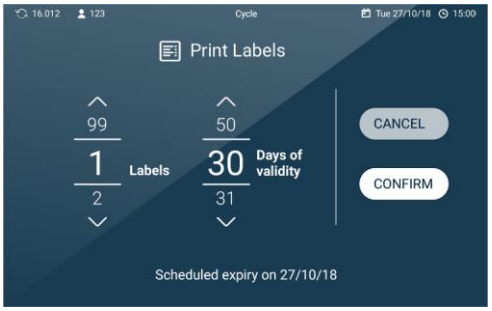
The choice between report and extended report determines also the report format available for automatic download or download via USB.



6.1.7.1. PRINT LABELS

If Labels option is selected, the dedicated menu is displayed when the door opens at the end of the cycle to allow setting:
- number of labels;
- package validity interval, in days.

The date that will be printed is indicated below.

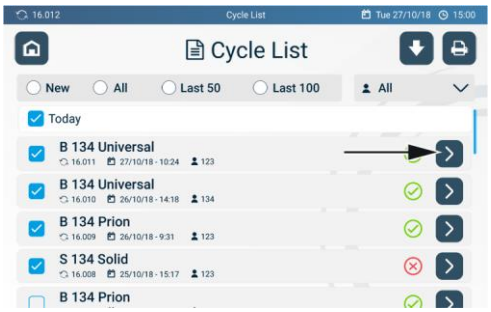


7. CYCLE LIST

From the Home page it is possible to access the CYCLE LIST menu in order to display, manage and download the cycle reports stored in the internal memory of the device.

Access CYCLE LIST to view the list of all the cycles performed by the device and the final outcome of each one of them.

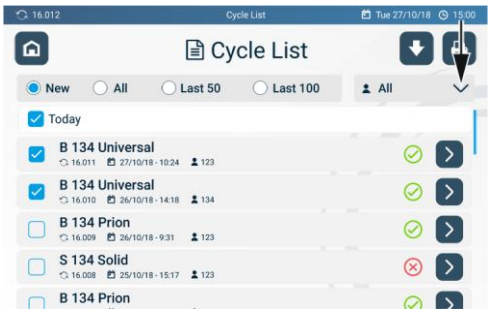
The indicated control allows displaying an individual cycle report on the screen



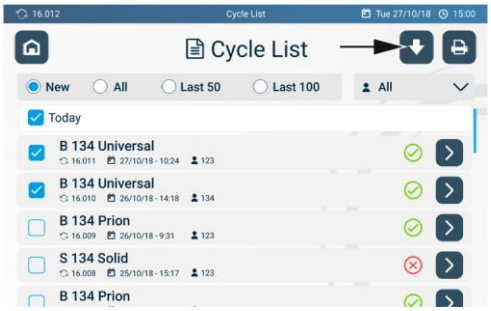
From CYCLE LIST it is possible to download the reports stored in the device on a USB memory support inserted in the front USB port.
The download can be customised by using the selection boxes on the left of the CYCLE LIST and/or by enabling one of the following filters:

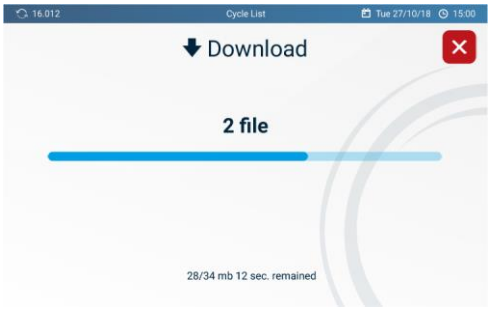
- NEW: to download reports never downloaded before.
- ALL: to download all cycle reports.
- LAST 50/100: to download the cycle reports corresponding to the selection made.

The OPERATOR control allows selecting the user based on which results are to be filtered by accessing a dedicated page.

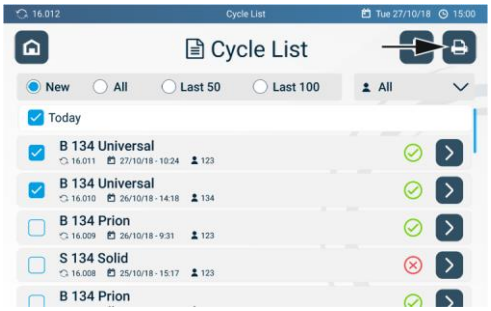


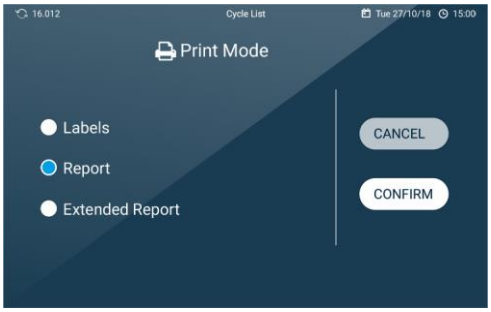
The control activates the download of reports, based on the filters enabled. Reports are downloaded in PDF format.





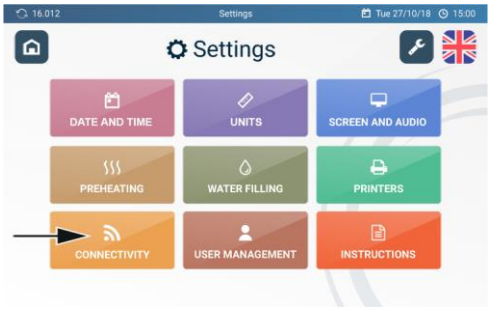
The control activates the printing of cycle reports, according to the mode selected.

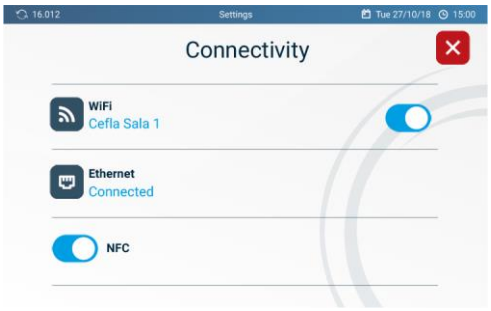




8. CONNECTIVITY

Select CONNECTIVITY to set WIFI, ETHERNET or NFC connection.



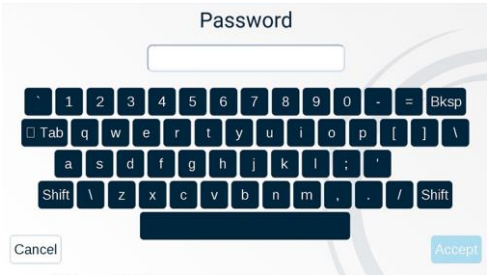


8.1. WIFI

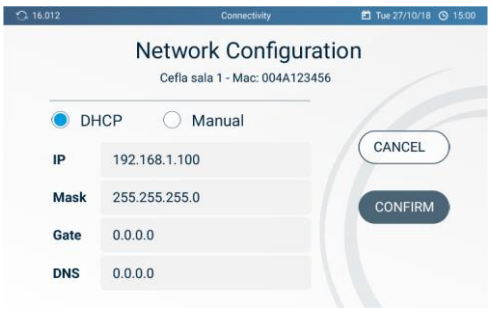
Use the switch to enable or disable Wi-Fi connection.

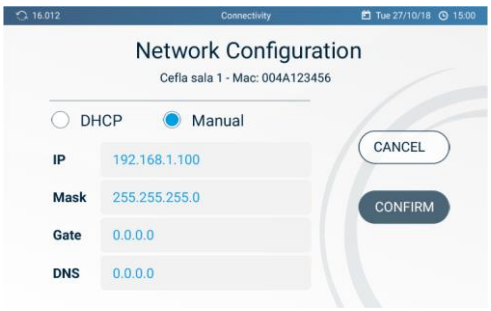
Select Wi-Fi to connect the sterilizer to a local Wi-Fi network.

When the desired network (SSID) has been selected, enter its access password (PSW).



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.



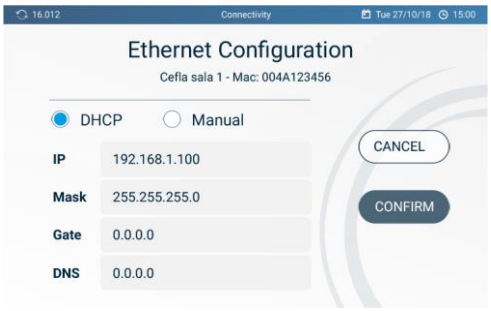


Make sure that Automatic DHCP configuration is selected.
With this selection all the numeric fields present on the screen are disabled (they get grey).
With this setting at each start the sterilizer requests the network DHCP server its configuration using the DHCP protocol.
According to DHCP server configuration the numbering received may change at each start.
The TCP-IP number assigned to the sterilizer appears in the Ethernet or Wi-Fi setting screen.
It is usually possible to set DHCP server in order to always assign the same IP number to a given device or to assign the same number to a given device for a predetermined period of time.
For these settings refer to the instruction manuals of your DHCP Server or of the local network Internet router.
These settings may require the “MAC address” of the sterilizer, for this information it is necessary to contact the technical service.

8.2. ETHERNET CONNECTIVITY

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

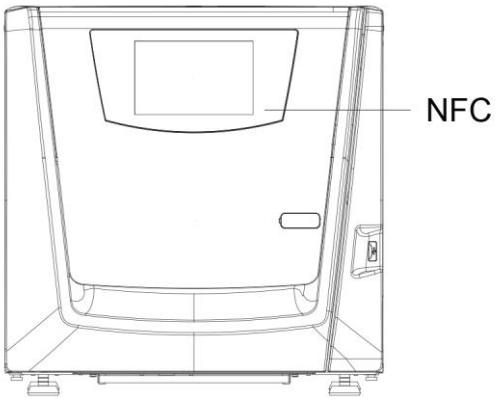
In this case too, it is possible to set DHCP as automatic or manual. In automatic mode the network configuration parameters are assigned automatically, in manual mode the network configuration parameters must be set manually.



Make sure that Automatic DHCP configuration is selected.
With this selection all the numeric fields present on the screen are disabled (they get grey).
With this setting at each start the sterilizer requests the network DHCP server its configuration using the DHCP protocol. According to DHCP server configuration the numbering received may change at each start.
The TCP-IP number assigned to the sterilizer appears in the Ethernet or Wi-Fi setting screen.
It is usually possible to set DHCP server in order to always assign the same IP number to a given device or to assign the same number to a given device for a predetermined period of time.
For these settings refer to the instruction manuals of your DHCP Server or of the local network Internet router.
These settings may require the “MAC address” of the sterilizer, for this information it is necessary to contact the technical service.

8.3. NFC

The control allows activating the NFC magnetic card reader located on the front display.



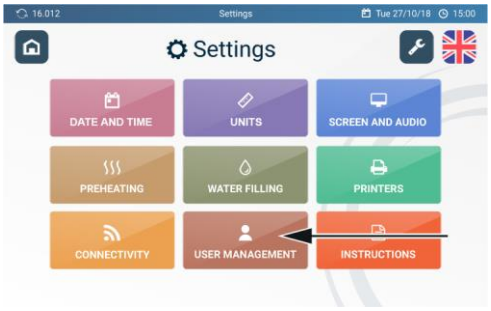
9. USER MANAGEMENT

Select USER MANAGEMENT to set the users.
Upon the first use, create the ADMIN user following the indications in the figure below.

The first user entered is given administrator rights.

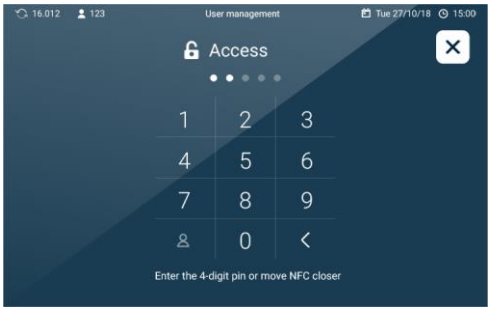
*When the user enters an incorrect PIN for 3 times, it is necessary to perform the unlocking procedure described **IN THE APPENDIX - USER PIN RESET**.*

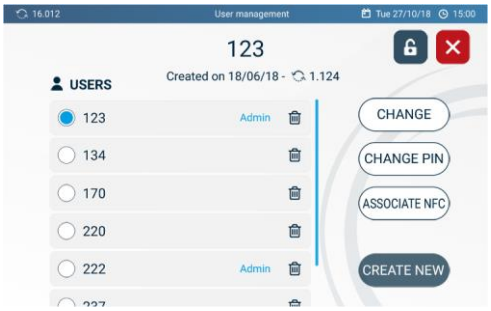
After you have entered the PIN, you can access the reserved administrator menu.
The ADMIN user can decide whether the sterilizer will request the PIN to the generic user at the cycle start (PIN Start) and/or at the cycle end (PIN End).



By accessing his/her own profile, the ADMIN user can:

- CREATE NEW - Create new users
- CHANGE - Associate users with specific sterilization cycles
- ASSOCIATE NFC - Associate users to NFC cards
- CHANGE PIN - Change personal identification PIN
- Manage user identification at the cycle start/end.
- Delete users.



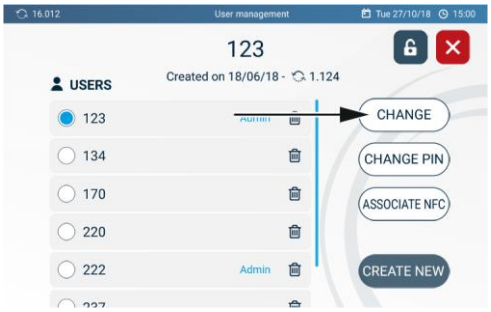


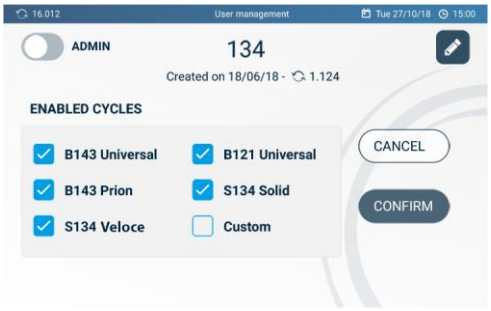
9.1. CREATE NEW

This function allows adding new users by entering a name and an identification PIN for each one of them.
It is possible to enter up to 30 users in the list.

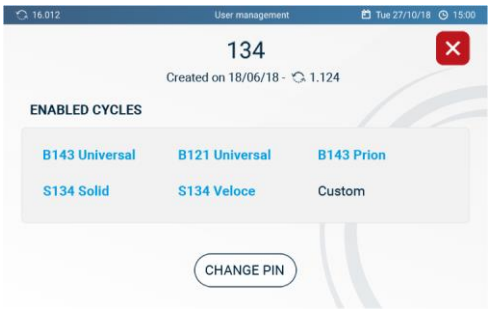
9.2. CHANGE

This function, reserved to ADMIN, allows associating one or more sterilization cycles with each user, choosing them among the six available ones.



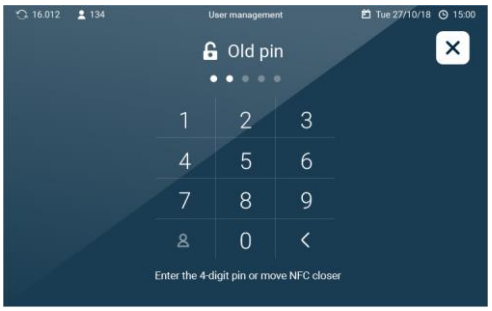


Each user will be able to check which are the cycles for which he/she is enabled by accessing his/her own profile.



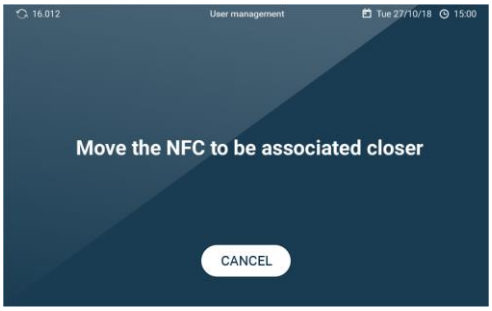
9.3. CHANGE PIN

This function allows each single user to change the identification PIN originally entered.



9.4. ASSOCIATE NFC

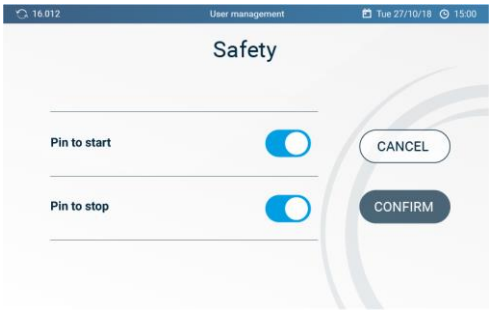
This function, reserved to ADMIN, allows associating an NFC card with each user.
To perform user/NFC card pairing, first activate the NFC magnetic card reader from SETTINGS.



9.5. USER PIN REQUEST

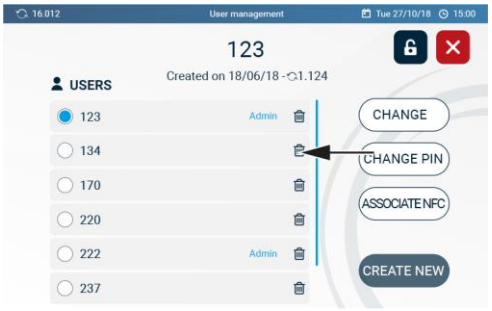
This function, reserved to ADMIN, allows activating the user identification request:
- Only at CYCLE START;
- Only at CYCLE END;
- At CYCLE START and END.

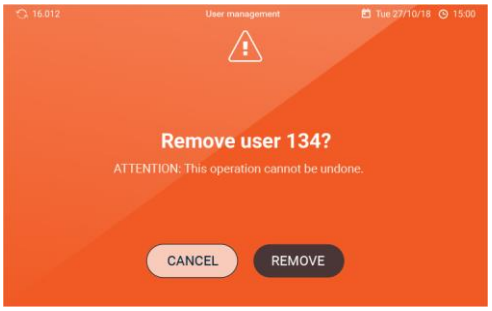
If the PIN is requested at CYCLE START and END, the system accepts the identification of two users for the two operations.
The cycle report, both in digital and printed format, will include the names of both users.
If the PRINT LABELS option is selected, labels will include the name of the user identified at the CYCLE END.

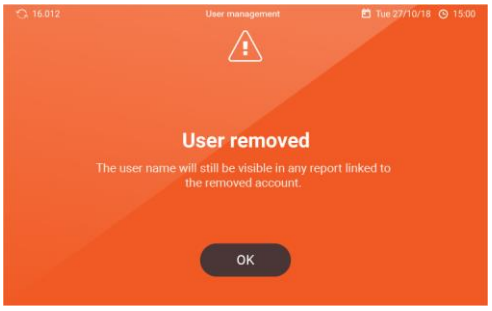


9.6. USER DELETION

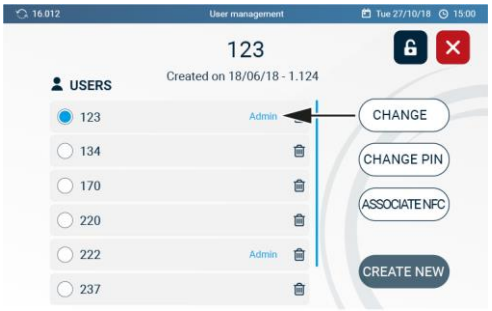
This function, reserved to ADMIN, allows eliminating the user entered in the list.
The user deletion is not reversible.
The name of the deleted user will still be shown in any cycle reports associated with the removed account.



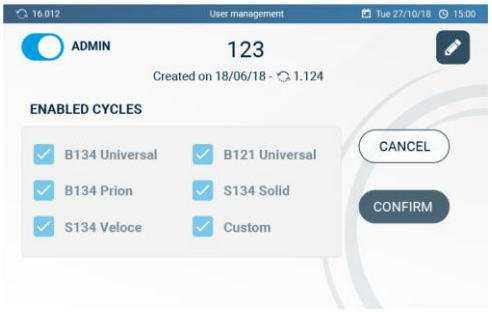




Select ADMIN user to view the summary data on the user.

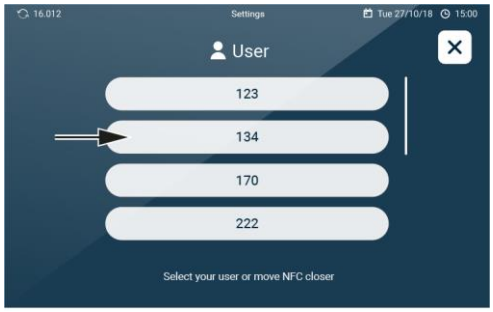


The ADMIN user can choose which cycles the selected user is authorised to perform by pressing the corresponding icons.

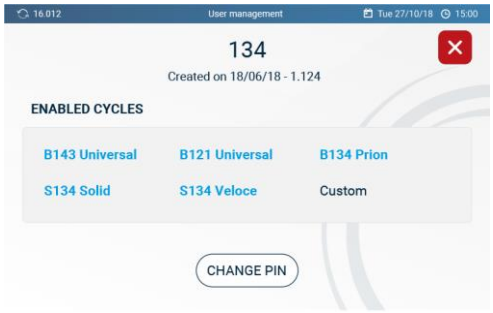


9.7. USERS LIST

Select the desired user and identify yourself with your personal PIN.
You will access the screen containing data of the selected user.

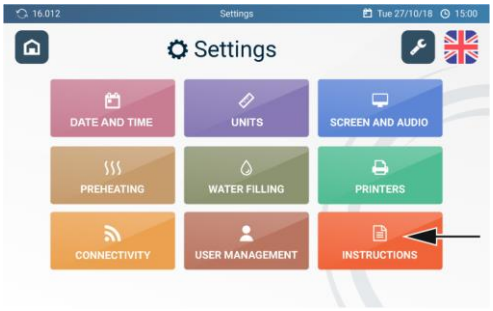


Once entered, a NON ADMIN user can see only a summary of his/her data, or change his/her PIN (see entering PIN - the following is requested in sequence: current PIN, new PIN, new PIN confirmation).

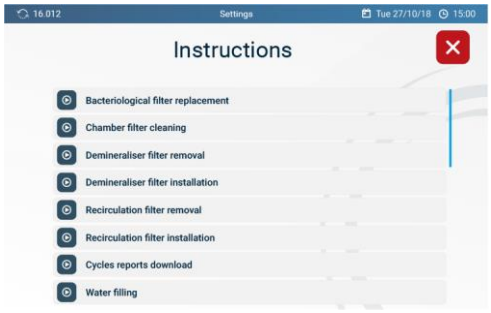


10. INSTRUCTIONS

Select INSTRUCTIONS to access the instructions and video tutorials for the user.



Press on the video of your interest to show the selected tutorial on the sterilizer display.



11. SERVICE

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

12. 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.

12.1. 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.

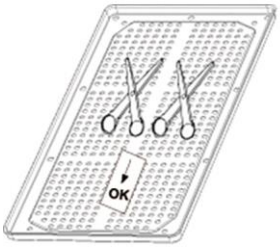
12.2. ARRANGING THE LOAD

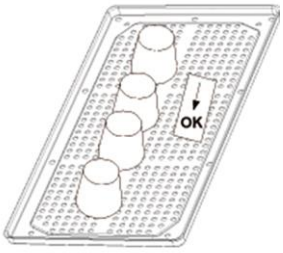
Always use personal protective equipment.

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.

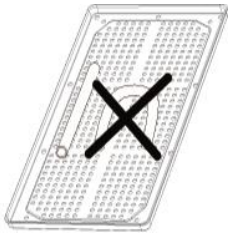




Place one sterilization chemical indicator per tray to indicate when the process is complete: this will allow avoiding an unnecessary repetition of the process on the same load or, worse, the use of unsterilised material. If packed material is sterilized, place the indicator inside one of the packages.

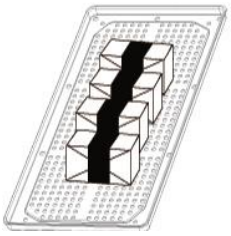
Note for rubber and plastic hoses:

- 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.

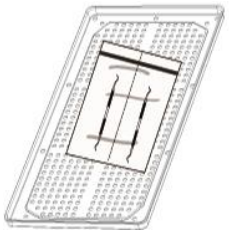


Notes for packages:

- 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 tray;
- 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.

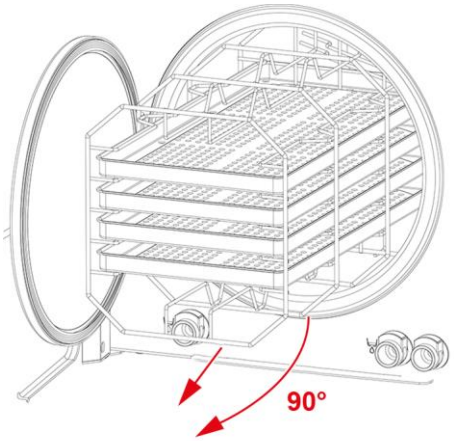
Since all instruments, or material in general, have a different structure, consistency and properties, the **most suitable program must be identified**, both to preserve the physical characteristics (avoiding or, in any case, limiting its alterations) and to ensure the best effectiveness of the sterilization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

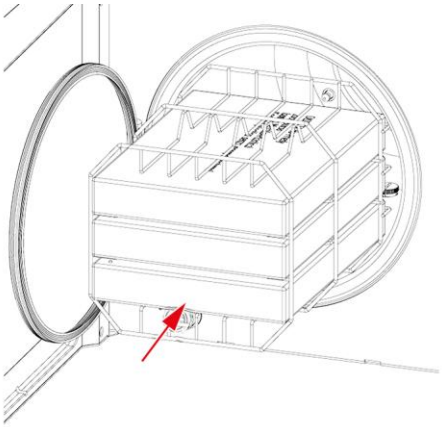
12.3. POSITIONING AND USE OF TRAY HOLDER SUPPORT

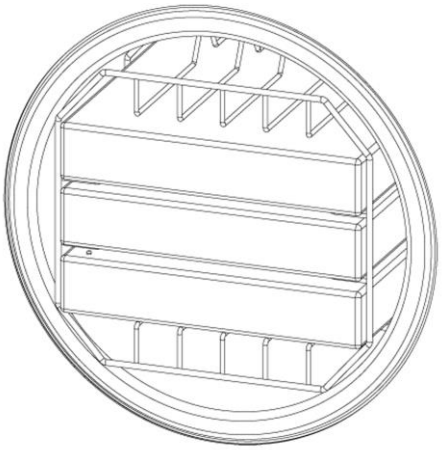
Tray holder support can be used in "tray" version (5/6 compartments based on the sterilizer model).

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.







13. 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.

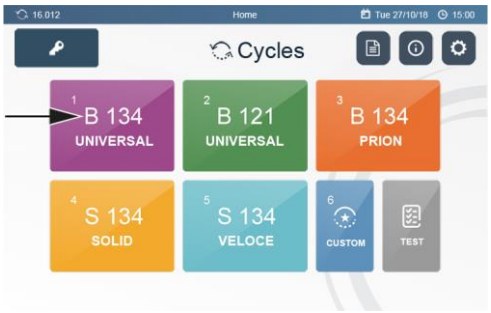
Available cycles are:
- **134 Universal B**
- **121 Universal B**
- **134 Prion B**
- **134 Solid S**
- **S 134 Veloce**
- **Custom (User-defined)**

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.

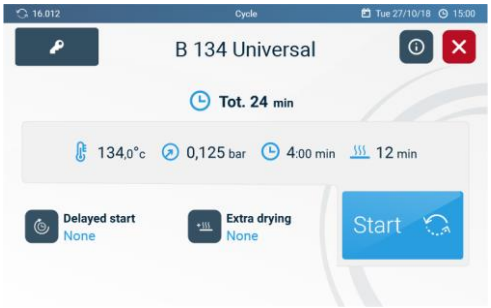
With this type of control, when you select a suitable sterilization program, you are guaranteed an effective sterilization under any conditions.

After inserting the load in the sterilization chamber (taking the precautions described in the section **"Preparing the material to be sterilized"**), select the desired sterilization cycle as follows:

To select the cycle, press the corresponding icon.

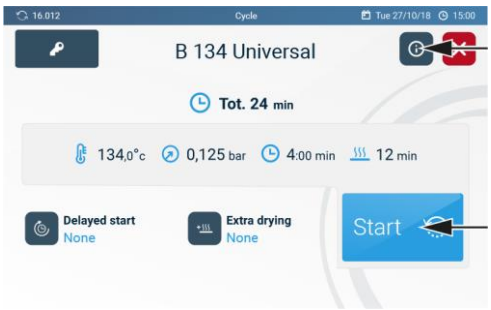


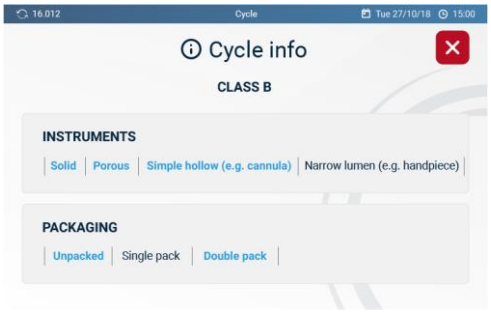
The cycle counter appears in the upper left corner.
Day and time are on the right.
The screen shows all the summary information of the selected cycle:
- Cycle name;
- Total cycle time;
- Process rated temperature;
- Process rated pressure;
- Process time;
- Any set extra drying;
- Any set delayed start.



Press INFO to display the information on the type of load compatible with the selected cycle.

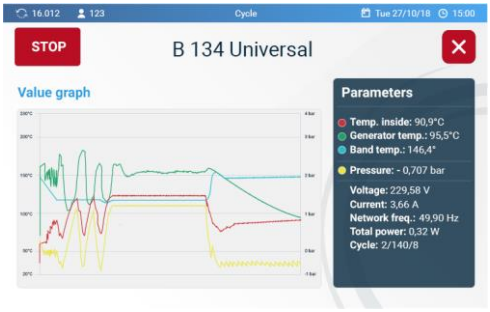
Press START to start the cycle.





When the cycle is in progress, the following screen is displayed:

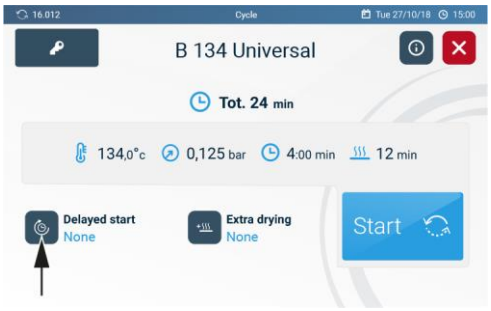
Press INFO to display the information on the cycle parameters and the graph.

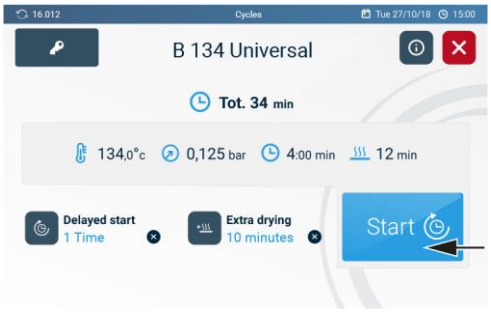


13.1. DELAYED START

The control allows accessing the start delay setting, expressed in hours.
The delayed start date and time are indicated at the centre.
The cycle start screen and the set delay value will be displayed when confirming
Delayed start can be activated/disabled without changing the setting.

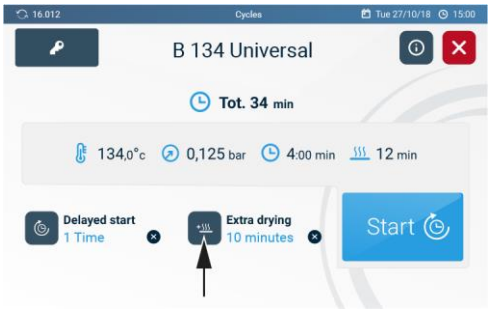
Press Delayed start to set the desired delay time for the start of the selected cycle.





13.2. EXTRA DRYING

The control allows accessing the additional drying time setting, expressed in minutes. The total drying time
is indicated at the centre.
The cycle start screen and the set extra drying time value will be displayed when confirming.
Extra drying can be enabled/disabled without changing the setting.



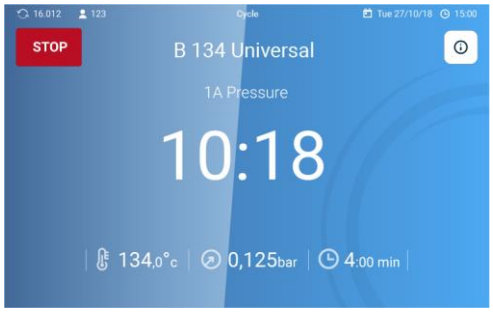
Set the minutes of drying you want to add to the standard drying time.

Confirm by pressing CONFIRM.

Upon the next use, just press the extra drying key to activate the values previously set.

13.3. EXECUTION OF THE CYCLE

Taking as example the most complete and significant sterilization cycle, i.e. the **B 134°C UNIVERSAL** program, characterised by fractionated pre-vacuum, the cycle sequence is as follows:

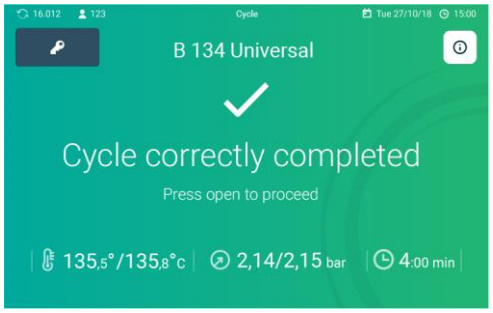


WARMING UP
FIRST VACUUM PHASE
FIRST PRESSURE RISE
SECOND VACUUM PHASE
SECOND PRESSURE RISE
THIRD VACUUM PHASE
THIRD PRESSURE RISE
STERILIZATION
STEAM DISCHARGE
DRYING
VENTILATION
CYCLE COMPLETION

13.4. CYCLE OUTCOME

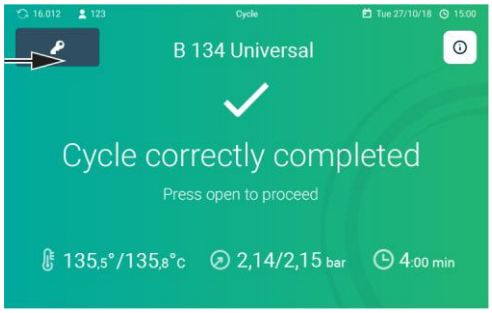
At the end of the cycle it is important to check the sterilization process outcome.
If the message **"CYCLE CORRECTLY COMPLETED"** is displayed, it means that the cycle has been correctly completed without any interruption due to any type of alarms and that the **complete asepsis** of material is ensured.
At the end of the cycle, the relevant screen will be displayed including indications on:

- Minimum/maximum temperature during the process;
- Minimum/maximum pressure during the process;
- Process time.



13.5. DOOR OPENING AT CYCLE END

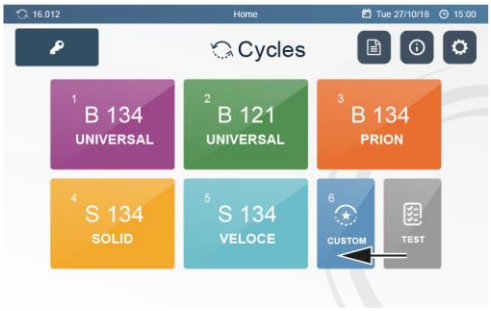
To open the sterilizer door, press the icon indicated in the figure:



User identification is required in this phase, if enabled.
If enabled, the label printing management screen will be displayed once the door has been opened.

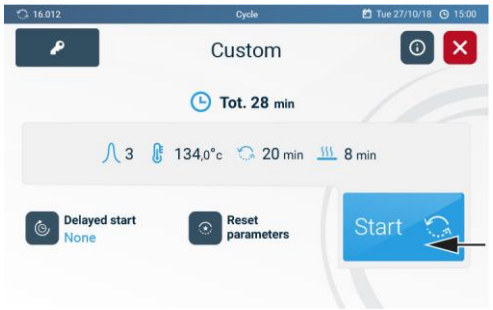
13.6. USER-DEFINED CYCLE

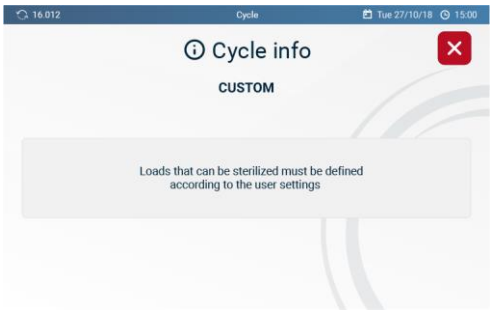
To set parameters press CUSTOM



Select the type of pre-vacuum, the process temperature, the exposure time and extra drying if necessary.
Once selections have been made, save the settings with CONFIRM key.

Press INFO to display the information on the cycle parameters.
Press START to start the user-defined cycle.





14. 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.

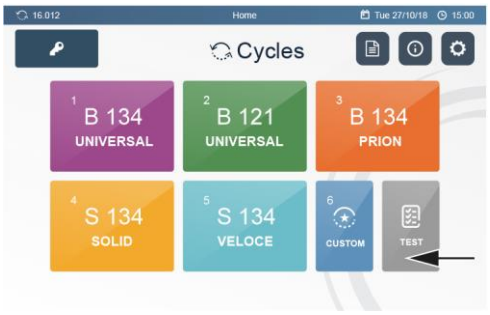
Such storage times may vary from country to country, according to the local legal requirements.

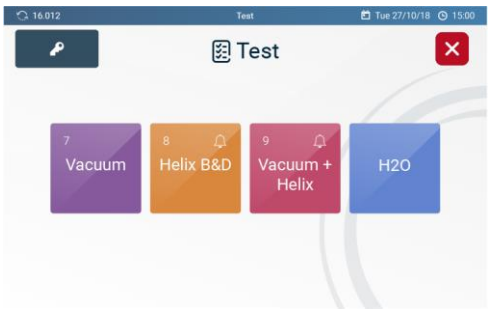
15. TEST PROGRAMS

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:

- **VACUUM TEST**;
- **HELIX** / **Bowie&Dick TEST**;
- There is also a program that performs the two combined tests **VACUUM + HELIX** / **Bowie&Dick TEST**;
- There is also a further test to check the water quality: **H2O TEST**.

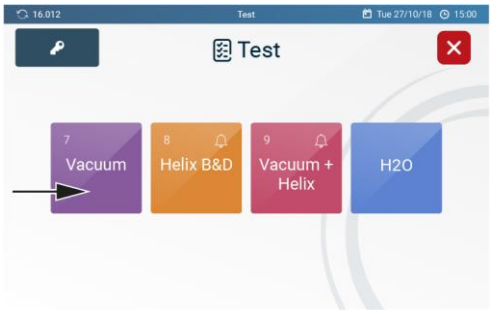


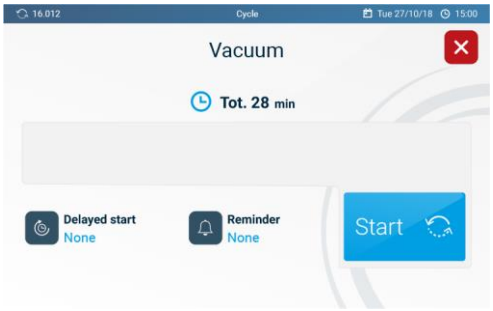


15.1. VACUUM TEST CYCLE

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 VACUUM TEST cycle, press the corresponding icon.





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.

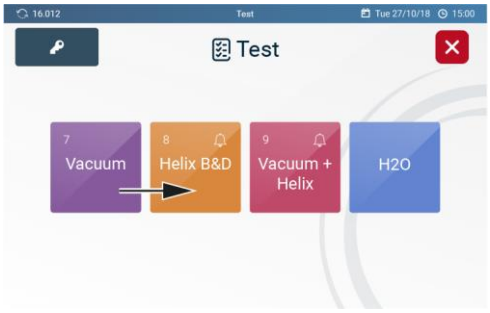
A high chamber temperature affects the variation in the vacuum value measured during the test; the system is therefore programmed to prevent execution of the test when the operating conditions are inadequate.
Close the door and start the program.
The vacuum phase starts immediately and the pressure value (bar) and the countdown from the start of the test cycle are shown on the display.

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.

15.2. HELIX / BOWIE DICK TEST

HELIX/Bowie&Dick TEST is a cycle at 134°C characterised by a sterilization phase having a special duration time (3.5 minutes); the cycle includes 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 **Helix/Bowie&Dick** cycle, press the corresponding icon.



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.

The test device and the chemical indicators to execute the helix/b&d test cycle are not provided with the device.
For information in this regard, contact technical service department (see appendix).

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.

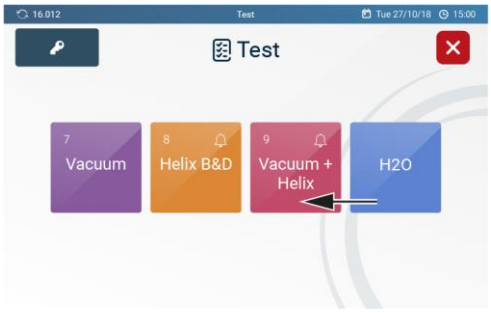
The test cycle takes place with a succession of phases similar to those described for a normal sterilization cycle.
At the end of the cycle, remove the test device from the chamber, open the cap and remove the indicator from its housing.
If the steam has correctly penetrated, the ink will have completely 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.

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.

The Bowie & Dick test package includes a chemical indicator inserted between various layers of porous material, consisting of a sheet with a special reagent ink on its surface.
If the steam penetration through the porous material layers is correct, it will make the ink colour change evenly throughout the entire surface of the B&D indicator.
If this is not the case, the toning will not occur or will be partial.

15.3. VACUUM CYCLE + HELIX / BOWIE - DICK TEST

Select this option to perform a VACUUM TEST cycle and a HELIX / Bowie&Dick TEST cycle in sequence.

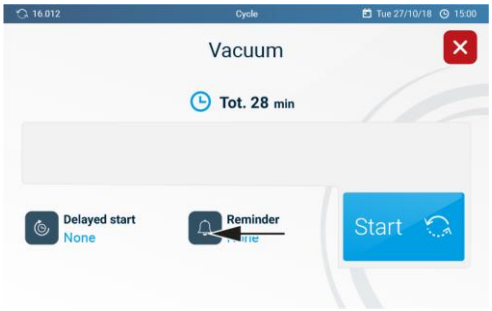


To this end, place the test device on the central tray without inserting other material.
Close the door and start the cycle.
The program will execute the two cycles in succession.
Check the results as described in the previous paragraphs.

The presence of the Helix test device does not alter the performance and the outcome of the vacuum test cycle.

15.4. TEST REMINDER

This function allows displaying, based on the chosen interval, a message recalling the performance of the relevant test.

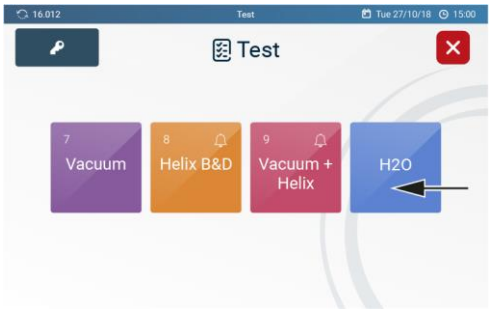


Set whether and when Test reminders are to be activated (Vacuum - Helix/Bowie&Dick Test- Vacuum + Helix/Bowie&Dick Test) based on the available options.
Once the fields have been set, activate the reminder with CONFIRM.
Reminders activate at 8 a.m. of the selected day or upon switching-on (if it occurs 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).

15.5. H2O TEST

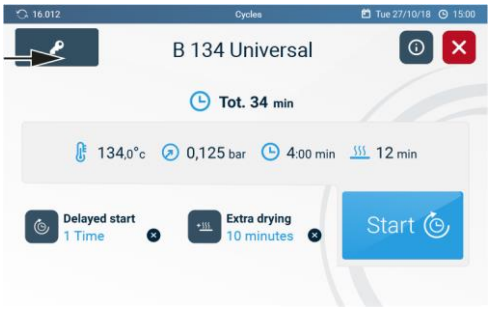
Select this option to test the water quality.



The water conductivity is automatically measured at each sterilization or test cycle start.

15.6. DOOR OPENING

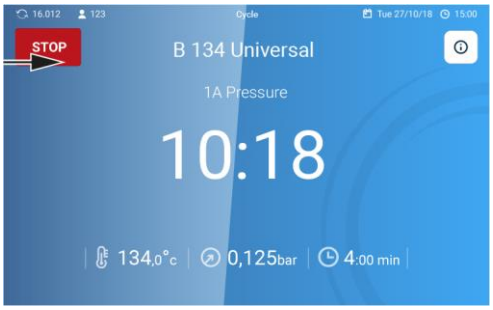
Press the indicated icon to open the sterilizer door when cycle ends.

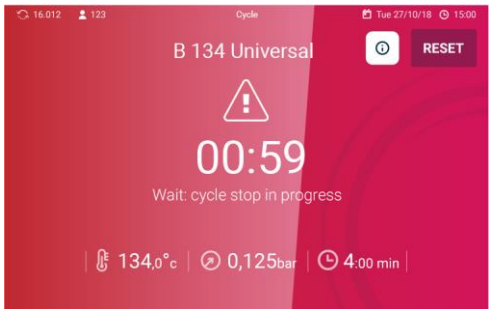


The door opens and remains ajar.
Then the door can be opened manually.

15.7. MANUAL INTERRUPTION

The cycle can be manually interrupted by the operator at any time, by keeping the icon shown in the figure **pressed for about three seconds**.



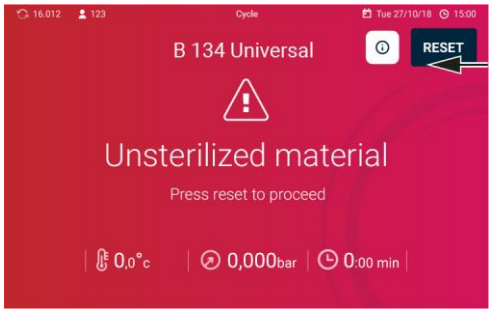


The command generates **E999 error** since the cycle could not finish correctly.

In certain phases of the cycle, the manual interruption activates an automatic cleaning procedure of the internal hydraulic circuit, that can last some minutes.
For a complete description of the alarms, refer to the appendix "Alarms".

In certain phases of the cycle, the manual interruption activates an automatic cleaning procedure of the internal hydraulic circuit, that can last some minutes.
For a complete description of the alarms, refer to the appendix "Alarms".

Press RESET for 3 seconds to open the door.



16. USED WATER DRAIN

This operation is required only if a sterilizer without used water recirculation filter is used.

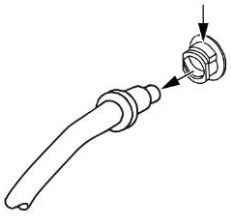
When the water maximum level is reached, a specific message is displayed.

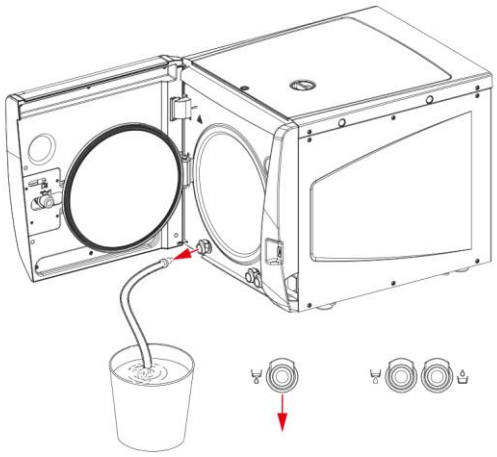
Open the door and continue as follows:

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.

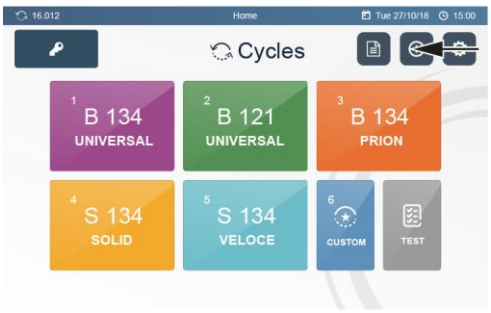
Detaching the tube

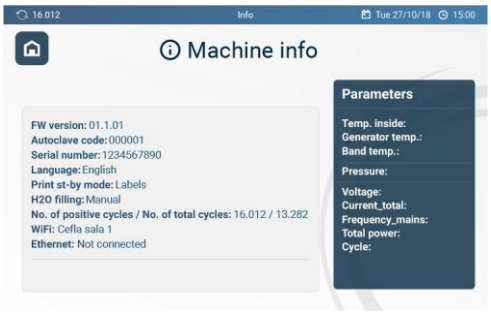




17. STERILIZER INFORMATION

Press INFO to display the summary of the sterilizer information.





18. DATA MANAGEMENT

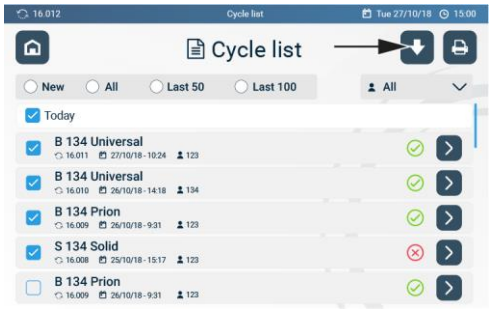
To access DATA MANAGEMENT section press the relevant icon.

See also the chapter CYCLE LIST.

18.1. CYCLE DOWNLOAD

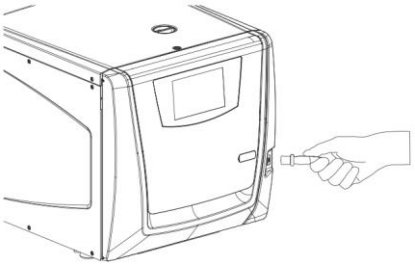
Before carrying out the following operations insert the USB key.
It is possible to copy data about the cycles carried out, stored in the inner memory of the sterilizer, on a USB key.

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

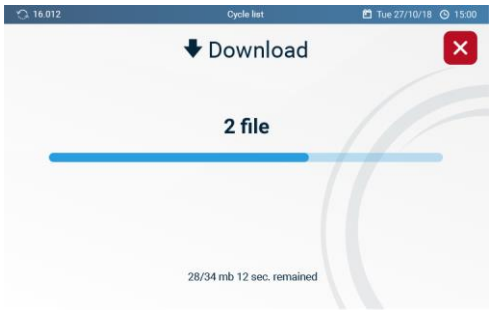


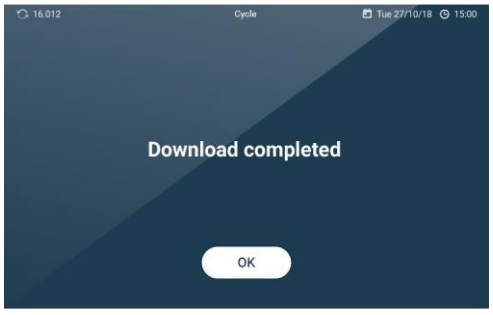
USB key has to be formatted according to the indications included in: Appendix - summary table technical characteristics.

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



The sterilization / test cycle report files are in PDF format.





19. 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.

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

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C UNIVERSAL	134	2.1	4(*)	B	F	13	40	550	0.75	Unwrapped porous materials	1.00	0.30	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	0.75	0.25	0.25	
										Porous materials in double pack	0.60	0.20	0.20	
										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	
134°C PRION	134	2.1	18	B	F	13	54	600	0.85	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	
										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	
121°C UNIVERSAL	121	1.1	20	B	F	13	56	600	0.75	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	
										Solid and hollow materials in single pack	3.00	1.00	0.50	
										Unwrapped solid and hollow materials	6.00	1.20	0.25	

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Solid and hollow instruments in double pack	1.50	0.50	0.25	
134°C VELOCE	134	2.1	4(*)	S	F	1	22	450	0.65	Unwrapped hollow instruments	2.00	1.20	0.50	
										Unwrapped solid instruments	2.00	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	-	-	-	-	-	-	

19.2. SUMMARY TABLE OF 17 120 V CYCLES

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
135°C HOLLOW WRAPPED	135	2.2	4(*)	B	F	13	48	550	0.75	Unwrapped porous materials	1.00	0.30	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	0.75	0.25	0.25	
										Porous materials in double pack	0.60	0.20	0.20	
										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	
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	
121°C RUBBER & PLASTIC	121	1.1	20	B	F	13	67	600	0.75	Unwrapped porous materials	1.00	0.30	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	0.75	0.25	0.25	
										Porous materials in double pack	0.60	0.20	0.20	
										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	
135° HOLLOW UNWRAPPED	135	2.2	4(*)	S	F	4	40	550	0.65	Unwrapped hollow instruments	6.00	1.20	0.50	
										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	

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	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	-	-	-	-	-	-	

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

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C UNIVERSAL	134	2.1	4(*)	B	F	15	44	700	0.8	Unwrapped porous materials	1.20	0.40	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	
										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	
134°C PRION	134	2.1	18	B	F	15	58	750	0.9	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	
										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	
121°C UNIVERSAL	121	1.1	20	B	F	15	61	750	0.8	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	
										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	
134°C VELOCE	134	2.1	4(*)	S	F	1	25	500	0.65	Unwrapped hollow instruments	2.00	1.50	0.50	

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped solid instruments	2.00	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	-	-	-	-	-	-	

19.4. SUMMARY TABLE OF 22 120 V CYCLES

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
135°C HOLLOW WRAPPED	135	2.2	4(*)	B	F	15	53	550	0.8	Unwrapped porous materials	1.20	0.40	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	
										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	
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	
121°C RUBBER & PLASTIC	121	1.1	20	B	F	15	73	750	0.8	Unwrapped porous materials	1.20	0.40	0.30	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.00	0.30	0.25	
										Porous materials in double pack	0.75	0.25	0.20	
										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	
135° HOLLOW UNWRAPPED	135	2.2	4(*)	S	F	5	44	750	0.7	Unwrapped hollow instruments	7.50	1.50	0.50	
										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
										Unwrapped solid and hollow materials "B"	7.50	1.20	0.50	

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	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	-	-	-	-	-	-	

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

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
134°C UNIVERSAL	134	2.1	4(*)	B	F	17	54	900	0.8	Unwrapped porous materials	1.50	0.50	0.50	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.25	0.35	0.35	
										Porous materials in double pack	0.90	0.30	0.30	
										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	
134°C PRION	134	2.1	18	B	F	17	68	950	1	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	
										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	
121°C UNIVERSAL	121	1.1	20	B	F	17	67	950	0.9	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	
										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	
134°C VELOCE	134	2.1	4(*)	S	F	1	28	600	0.65	Unwrapped hollow instruments	2.00	1.50	0.50	

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
										Unwrapped solid instruments	2.00	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	-	-	-	-	-	-	

19.6. SUMMARY TABLE OF 28 120 V CYCLES

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	MAX TOTAL MASS (kg)	MAX MASS PER TRAY (kg)	MAX MASS PER ITEM (kg)	
135°C HOLLOW WRAPPED	135	2.2	4(*)	B	F	17	65	900	0.8	Unwrapped porous materials	1.50	0.50	0.50	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.25	0.35	0.35	
										Porous materials in double pack	0.90	0.30	0.30	
										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	
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	
121°C RUBBER & PLASTIC	121	1.1	20	B	F	17	80	950	0.8	Unwrapped porous materials	1.50	0.50	0.50	For wrapped materials and instruments (single and double pack), it is advisable to use the 3-tray configuration
										Porous materials in single pack	1.25	0.35	0.35	
										Porous materials in double pack	0.90	0.30	0.30	
										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	
135° HOLLOW UNWRAPPED	135	2.2	4(*)	S	F	6	50	950	0.7	Unwrapped hollow instruments	9.00	1.50	0.50	
										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

CYCLE DESCRIPTION	NOMINAL VALUES				BASIC CYCLE PARAMETERS					STERILIZABLE MATERIALS				NOTES
	Temperature (°C)	Pressure (bar)	Retention time (min)	Cycle type (EN 13060:2014)	Pre-vacuum (F=fractionated; S=single)	Standard drying (min)	Total cycle time (Max filling)	Max H2O consumption (ml/cycle)	Average energy consumption (kWh/cycle)	TYPE	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 "hollow loads" in this manual refers both to "narrow-lumen" elements (section 3.18 EN 13060:2014) and "simple hollow" elements (section 3.30 EN 13060:2014).
The term "hollow load B" refers ONLY to "simple hollow" elements (section 3.30 EN 13060:2014).

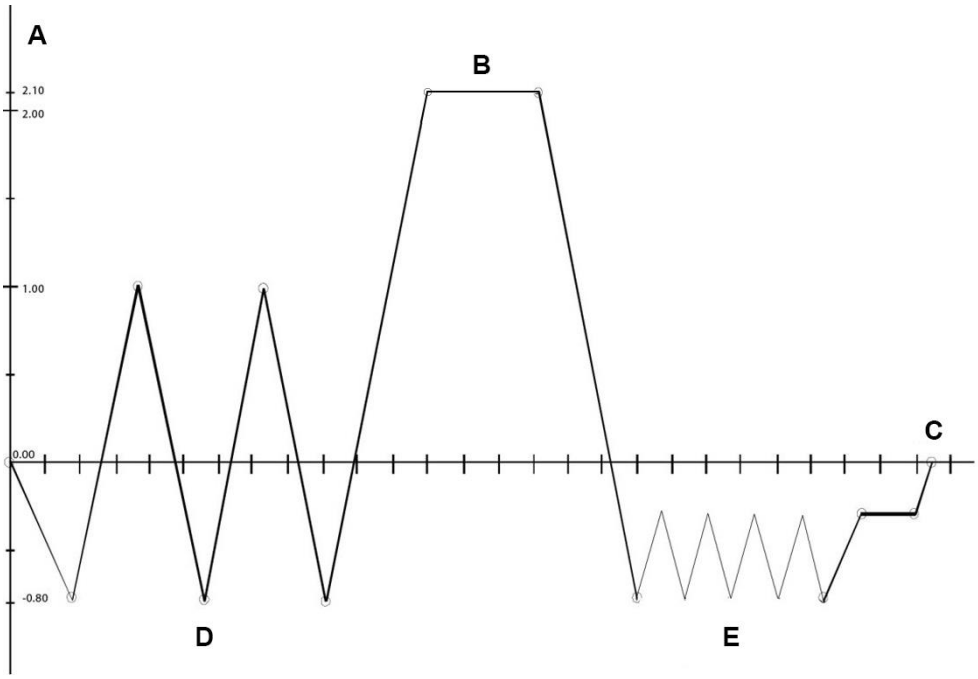
PRESSURE, TIME AND TEMPERATURE In compliance with EN 13060: 2014 for operating cycles						
134°C cycles						
EN 13060:2014		**Time (minutes)**	**Min temperature**	**Max temperature**	**Min pressure (bar)**	**Max pressure (bar)**
1	CS	---	---	---	---	---
t1	1PV	---	---	---	-0.81 * (-0.76)	-0.79 * (-0.74)
t2	1PP	---	---	---	+0.97 * (+0.27)	+1.03 * (+0.33)
t3	2PV	---	---	---	-0.81 * (-0.76)	-0.79 * (-0.74)
t4	2PP	---	---	---	+0.97 * (+0.27)	+1.03 * (+0.33)
t5	3PV	---	---	---	-0.81 * (-0.76)	-0.79 * (-0.74)
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 * (-0.76)	-0.79 * (-0.74)
t9	DE	---	---	---	---	---
2	CE	---	---	---	-0.02	+0.02
* Parameters referred to cycle S 134° VELOCE.						
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

19.7. STERILISATION PROGRAM DIAGRAM

PROGRAM
134°C UNIVERSAL
134°C – 4’ 00’’

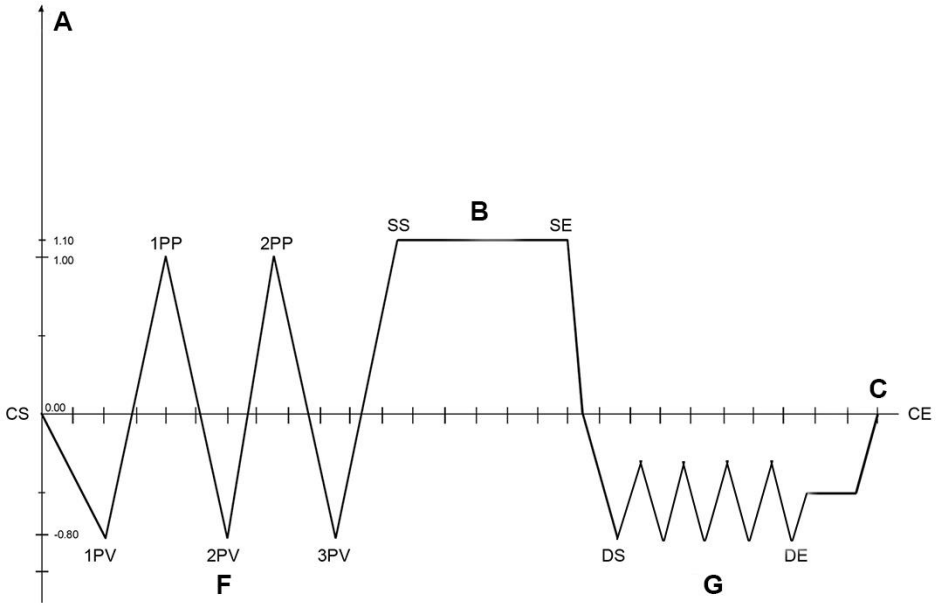
A PRESSURE (BAR)
B PROCESS
C TIME (MIN)
D FRACTIONATED VACUUM
E VACUUM DRYING

PROGRAM
134°C PRION
134°C – 18’ 00’’



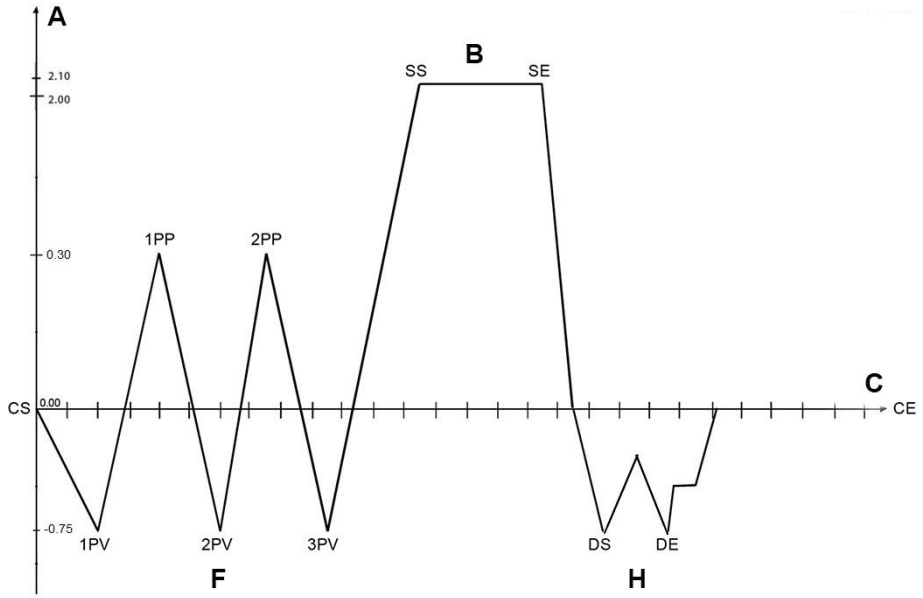
PROGRAM
121°C UNIVERSAL
121°C – 20’ 00’’

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



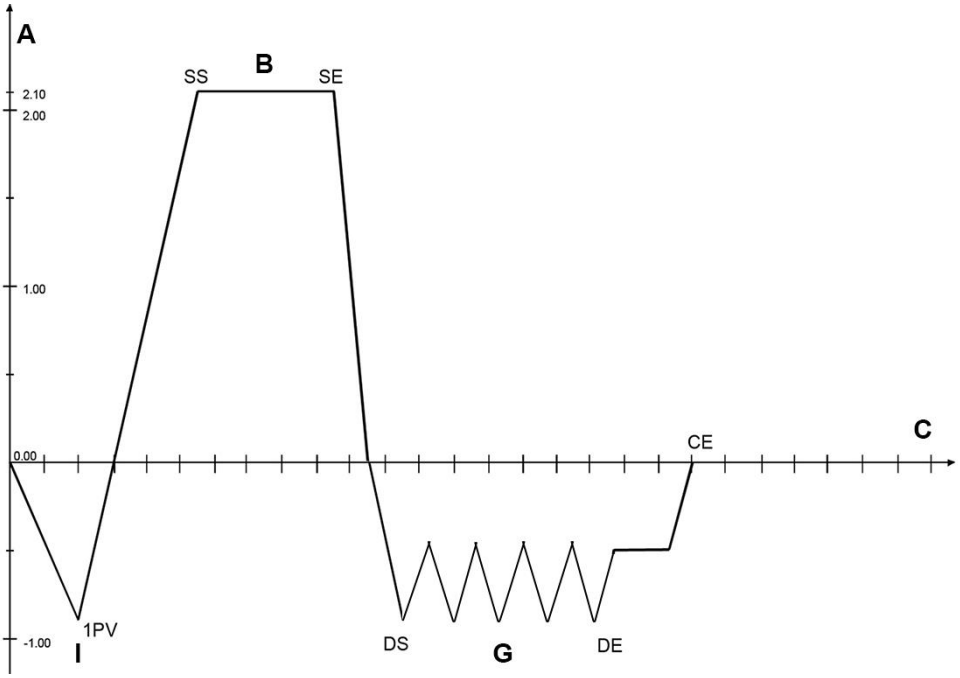
PROGRAM
134°C VELOCE
134°C – 4’00’’

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



PROGRAM
134°C SOLID WRAPPED
134°C – 4’00’’

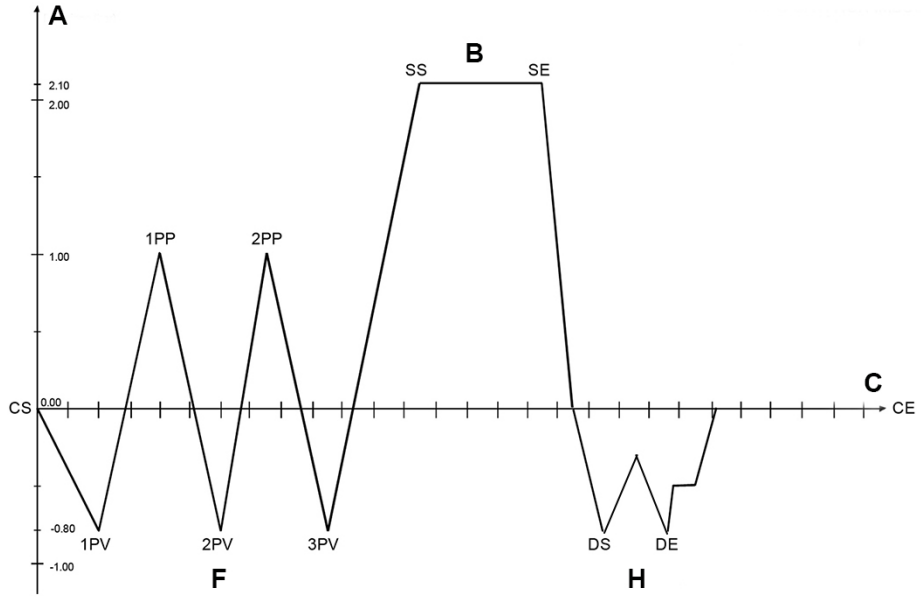
A PRESSURE (BAR)
B PROCESS
C TIME (MIN)
I SINGLE PRE-VACUUM
G LONG DRYING



19.8. DIAGRAMS OF THE TEST PROGRAMMES

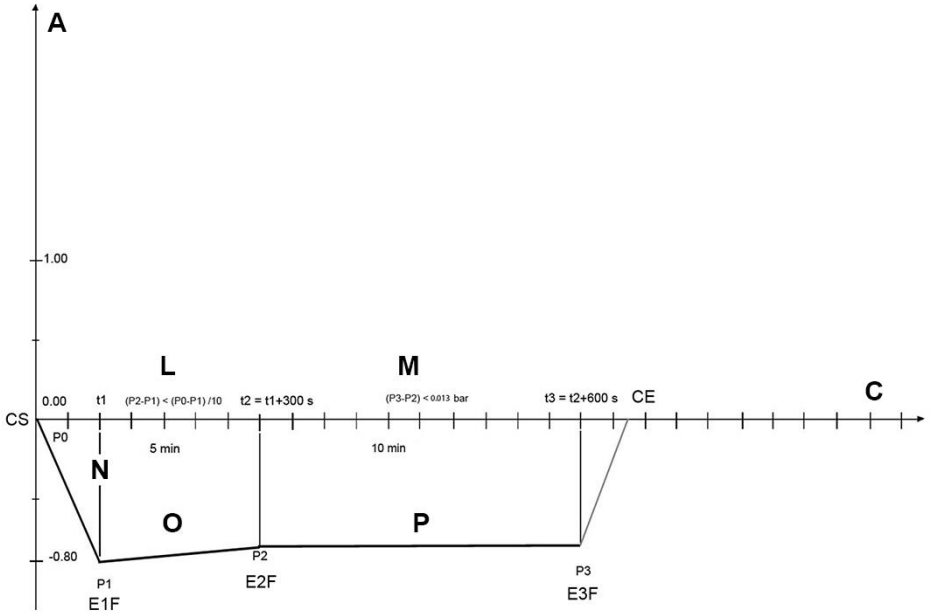
PROGRAM
HELIX B&D TEST
134°C – 3'0''

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



20. 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 **periodic sterilizer validation**, 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.

20.1. 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
UPON EVERY WATER FILLING OPERATION	Clean/disinfect filling/drainage tanks
UPON EVERY FILTER REPLACEMENT	
PERIODICALLY	See Scheduled Maintenance messages
YEARLY	Validate sterilizer ***(see scheduled maintenance)***

20.2. 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.
Press “REMIND” 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

TANK FILTER CLEANING

BOILER GASKET REPLACEMENT

GENERAL SERVICE

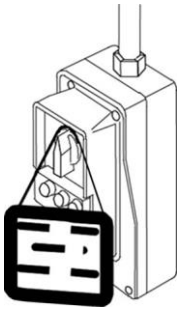
A regular maintenance is essential to achieve the best performance of the device.
Periodically, a message will be displayed requesting that the above maintenance operations are performed.
For further information or in case of doubt, contact the technical service: if they have performed regular maintenance 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.



20.3. DESCRIPTION OF MAINTENANCE INTERVENTIONS

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

20.3.1. 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.

20.3.2. 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).

20.3.3. 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.

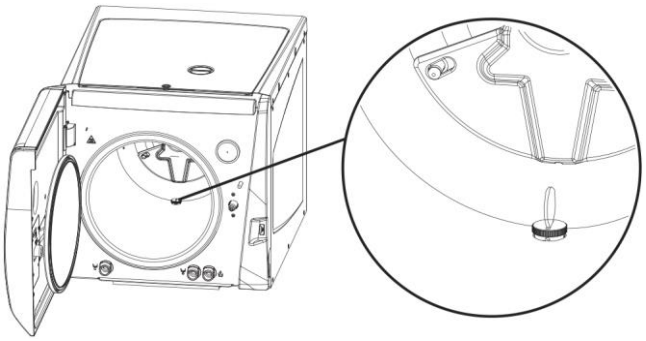
20.3.4. BOILER FILTER CLEANING

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 <u>impossible</u> to recover the filter, replace it with a new one.
Refit everything operating <u>in reverse order</u> and **<u>making sure</u>** to screw the union in such a way that the drain holes are positioned **at the level of the boiler wall**.

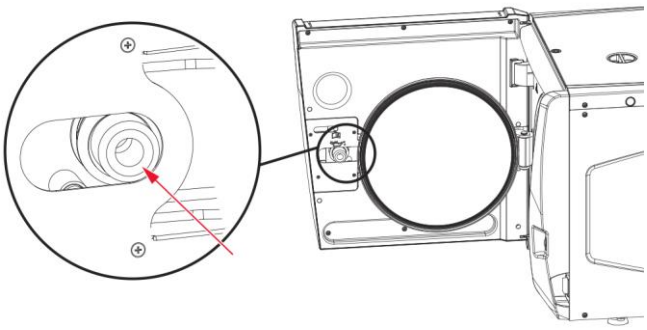
Properly fit the filter in its housing.
A partial fitting may damage the component.



20.3.5. 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).

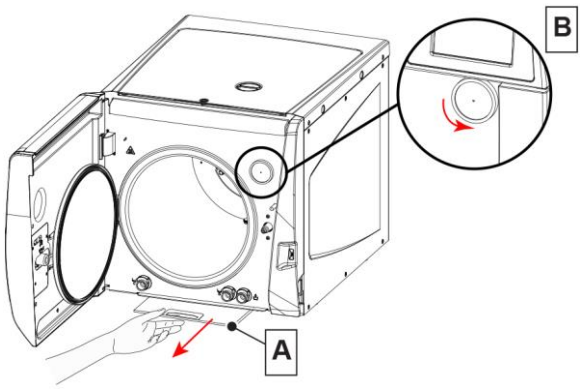
Wear single-use gloves before application.
Essentially, the lubricant is not irritant to the skin; nevertheless, it may cause unpleasant effects if it accidentally comes into contact with eyes.
In case of contact with eyes, rinse with plenty of water.



20.3.6. 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.



20.3.7. 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.

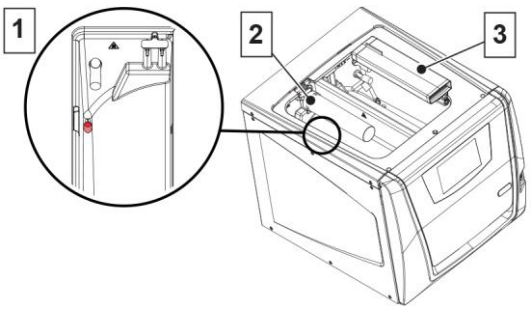
A spare bacteriological filter is provided with the device.
If you need spare parts of this component, refer to technical assistance appendix.

20.3.8. CLEANING AND DISINFECTION OF FILTERS AND WATER TANKS

Clean and disinfect filters (**1, 2, 3**) and the tank's internal walls only with a disposable towel/paper wipe soaked in 70%
ethyl alcohol.

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

Empty the tanks containing the sterilizer's filling and discharge water, remove any deposits around the filters (**1, 2, 3**) at the bottom of the tanks (see figure).
After extracting and cleaning the filters (**1, 2, 3**), thoroughly clean inside the tanks with a cloth soaked in 70% ethyl alcohol.
After cleaning the filters, refit them inside the tanks.



20.3.9. DEMINERALIZATION SYSTEM CARTRIDGE REPLACEMENT

The sterilizer is equipped with an integrated demineralisation system that performs an automatic check of the water conductivity value, including a set of signals.
If conductivity exceeds 20 µS/cm, the message E008 - “FILTER NEARLY EXHAUSTED” is displayed. It is possible to carry on using the device without any further action required.
If conductivity exceeds 60 µS/cm, the message E009 - “QUALITY H2O BAD / CHANGE FILTERS / CHANGE WATER” is displayed. The ion-exchange resin cartridge with must be changed.
Proceed as follows:
- Turn the sterilizer off;
- Remove the upper cover;
- Empty the filling tank through the union;
- Remove the cartridge by lifting the front part first, then the rear part;
- Replace the cartridge following the initial installation procedure (see INTEGRATED DEMINERALISATION SYSTEM).

20.3.10. RECIRCULATION SYSTEM CARTRIDGE REPLACEMENT

Replace the filter in case of alarms signalling the exhaustion of the recirculation filter or at the end of 500 machine’s cycles.
Proceed as follows:
- Turn the sterilizer off;
- Remove the upper cover;
- Empty the discharge tank through the union;
- Move the locking lever backward;
- Lift the filter;
- Replace the filter following the installation procedure indicated in the chapter “INSTALLING THE RECIRCULATION FILTER INSIDE THE TANK”.

Water may come out from the demineralisation system cartridge.

20.3.11. BOILER GASKET REPLACEMENT

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

20.4. 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.*

20.5. 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.

20.6. 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.

21. 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.

21.1. 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 sterilizer does not power-on.	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 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 resting surface.	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.
	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 humidity on the material and/or instruments at the end of the program.	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"**).
	Load not correctly positioned.	Position the load, in particular the wrapped one, as per the indications. (See **Chapter** "**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**").
Traces of oxidation or spots on instruments	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.

PROBLEM	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Quality of the distilled water not adequate.	Empty the tank and fill it with high-quality distilled water. (See Water supply characteristics in **"Technical characteristics" Appendix).**
	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).**

22. 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: **Exxx** **(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.
Code format: **Hxxx** **(xxx = identification number 000 ÷ 999)**

S = SYSTEM ERROR
Electronic system error (HW-FW).
Code format: Sxxx **(xxx = identification number 000 ÷ 999)**

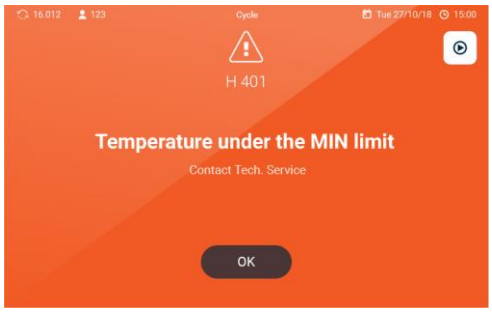
In case of alarm, switch off the device only after having followed the indications displayed and having carried out the reset (see "Resetting the system" paragraph).

22.1. 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**.

22.2. 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.



23. SYSTEM RESET

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

1 Pressing the OK button;

2 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 menu.

After the RESET, and any technical intervention necessary to remove the fault, the device will be ready to run a new program.

Never turn off the device before carrying out the reset.

24. ALARM CODES

The **list** of alarm codes, the relevant messages displayed and RESET modes, are indicated in the following table:

24.1. 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
E000	Black-out	POWER OUTAGE CONTACT TECH. SERVICE	2, 3
E001	Voltage of power supply line too high	OVERVOLTAGE CONTACT TECH. SERVICE	2, 3
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 discharge 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 filling tank	QUALITY H2O BAD CHANGE FILTERS CHANGE WATER	3
E020	Door lock system (closing) activation time-out exceeded	DOOR CLOSING ERROR CONTACT TECH. SERVICE	2, 3
E021	Door lock system (opening) activation time-out exceeded	DOOR OPENING ERROR CONTACT TECH. SERVICE	1
E022	Door lock microswitches failure.	DOOR LOCK PROBLEM CONTACT TECH. SERVICE	1
E030	The water in the feed tank is at minimum level (MIN)	LOAD TANK MINIMUM LEVEL FILL TANK	2
E031	Maximum level of water in the drainage tank (MAX)	DISCHARGE TANK MAXIMUM LEVEL EMPTY TANK	2
E042	The MAX water level in the filling tank has been reached	LOAD TANK MAXIMUM LEVEL	1
E050	Reminder to run Vacuum Test cycle	TEST REMINDER RUN VACUUM TEST	1
E051	Reminder to run Helix Test cycle	TEST REMINDER RUN HELIX TEST	1
E052	Reminder to run Vacuum + Helix Test combined 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
E070	Preheating activation with door open	PREHEATING ACTIVATED IT IS ADVISABLE TO CLOSE THE DOOR	1
E131	The selected user is not authorised to start or download the cycle	USER NOT ENABLED	1
E142	Console firmware version is not the correct one compared to Process firmware version. Malfunctions may occur in graphic interface management or data management to the outside (e.g. cycle reports, data on Cloud)	CONSOLE FW VERSION NOT CORRECT PLEASE UPDATE THE FW	1
E900	Vacuum test failed (during TEST PHASE)	TEST FAILED SECOND STEP CONTACT TECH. SERVICE	3
E901	Vacuum test failed (during STAND-BY PHASE)	TEST FAILED FIRST STEP CONTACT TECH. SERVICE	3
E902	Vacuum test failed (vacuum pulse time-out exceeded)	TEST FAILED VACUUM NOT ACHIEVED CONTACT TECH. SERVICE	3
E998	Remote maintenance activity in progress	REMOTE SERVICE ACTIVE	1
E999	Manually interrupting the cycle	MANUAL INTERRUPTION	3

1 = OK (warning)
2 = OK + Stopped cycle start
3 = Cycle failed + OK + RESET

24.2. ALARMS (CATEGORY A)

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
A032	Problem with the level sensor of the filling tank	FILL.WATER LEVEL 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	2, 3
A102	Temperature sensor PT2 broken (steam generator)	GENERATOR PROBE PT2 OPEN CIRCUIT CONTACT TECH. SERVICE	2, 3
A103	Temperature sensor PT3 broken (heating element)	HEATING BAND PROBE PT3 OPEN CIRCUIT CONTACT TECH. SERVICE	2, 3
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	2, 3
A112	Temperature sensor PT2 short-circuited (steam generator)	GENERATOR PROBE PT2 SHORT-CIRCUIT CONTACT TECH. SERVICE	2, 3
A113	Temperature sensor PT3 short-circuited (heating element)	HEATING BAND PROBE PT3 SHORT-CIRCUIT CONTACT TECH. SERVICE	2, 3
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	2, 3
A117	Door motor overcurrent error	DOOR MOTOR OVERCURRENT	2, 3
A120	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH. SERVICE	2, 3
A121	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH. SERVICE	2, 3
A122	Reference heating element acquisition chain fault	PROCESS BOARD ERROR CONTACT TECH. SERVICE	2, 3
A126	Connection error with Wi-Fi module	Wi-Fi MODULE ERROR CONTACT TECH. SERVICE	1
A127	Communication error between graphical interface and process board through Can	CAN ERROR CONTACT TECH. SERVICE	2, 3
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
A130	Connection error with RGB led bar	RGB LED BAR ERROR CONTACT TECH. SERVICE	1
A131	Solenoid valve 1 failed	SOLENOID VALVE 1 ERROR CONTACT TECH. SERVICE	2, 3
A132	Solenoid valve 2 failed	SOLENOID VALVE 2 ERROR CONTACT TECH. SERVICE	2, 3
A133	Solenoid valve 3 failed	SOLENOID VALVE 3 ERROR CONTACT TECH. SERVICE	2, 3

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
A134	Solenoid valve 4 failed	SOLENOID VALVE 4 ERROR CONTACT TECH. SERVICE	2, 3
A135	Solenoid valve 5 failed	SOLENOID VALVE 5 ERROR CONTACT TECH. SERVICE	2, 3
A136	Solenoid valve 6 failed	SOLENOID VALVE 6 ERROR CONTACT TECH. SERVICE	2, 3
A137	Solenoid valve 7 failed	SOLENOID VALVE 7 ERROR CONTACT TECH. SERVICE	2, 3
A145	Faulty current draw detected	FAULTY CURRENT DRAW CONTACT TECH. SERVICE	2, 3
A146	Solenoid valves control driver fault	SOLENOID VALVE DRIVER ERROR CONTACT TECH. SERVICE	2, 3
A147	Door motor control driver fault	DOOR MOTOR DRIVER ERROR CONTACT TECH. SERVICE	2, 3
A201	Heating not executed within time-out (steam generator)	STEAM GENERATOR RESISTOR OPEN CIRCUIT CONTACT TECH. SERVICE	3
A202	Heating not executed within time-out (tube bundle heating element)	HEATING BAND OPEN CIRCUIT CONTACT TECH. SERVICE	3
A250	1st pulse with vacuum not reached within time out	1PV TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A251	1st rise back up to atmospheric pressure not reached within the time-out	ATM1 UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A252	1st pressure pulse not reached within the time-out	1PP UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A253	2nd pulse with vacuum not reached within time out	2PV TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A254	2nd rise back up to atmospheric pressure not reached within the time-out	ATM2 UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A255	2nd pressure pulse not reached within the time-out	2PP UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A256	3rd pulse with vacuum not reached within time out	3PV TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A257	3rd rise back up to atmospheric pressure not reached within the time-out	ATM3 UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A258	3rd pressure pulse not reached within the time out	3PP UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A260	Chamber depressurization not reached within time out	ATM3 DOWNSTROKE TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A261	Chamber levelling not reached within time out	PRESSURE LEVELLING TIMEOUT CONTACT TECH. SERVICE	3
A262	Vacuum pulsation during drying not executed within time-out	PD PRESSURE UPSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A353	1st drop to atmospheric pressure not completed within the time-out	ATM1 DOWNSTROKE TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A356	2nd drop to atmospheric pressure not completed within the time-out	ATM2 DOWNSTROKE TIMEOUT CHECK LOAD CHECK CHAMBER FILTER	3
A360	Vacuum pulsation after maintenance step not executed within time-out	SPD PRESSURE DOWNSTROKE TIMEOUT CONTACT TECH. SERVICE	3
A362	Chamber depressurisation during drying not reached within time-out	PD PRESSURE DOWNSTROKE TIMEOUT CONTACT TECH. SERVICE	3

1 = OK (warning)
2 = OK + Stopped cycle start
3 = Cycle failed + OK + RESET

24.3. 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	2, 3
H160	MPX pressure sensor short-circuited	PRESSURE SENSOR SHORT-CIRCUIT CONTACT TECH.SERVICE	2, 3
H400	Pconv/T ratio not balanced (Pconv>T) (STERILIZATION phase)	INCORRECT P/T RATIO CHECK LOAD	3
H401	T/Pconv ratio not balanced (T>Pconv) (STERILIZATION phase)	INCORRECT T/P RATIO CHECK LOAD	3
H402	Temperature over the MAX limit (STERILIZATION phase)	TEMPERATURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	3
H403	Temperature below the MIN limit (STERILIZATION phase)	TEMPERATURE BELOW MINIMUM LIMIT CONTACT TECH. SERVICE	3
H404	Floating temperature over the limit (STERILIZATION phase)	ERRATIC TEMPERATURE CONTACT TECH. SERVICE	3
H405	Pressure over the MAX limit (STERILIZATION phase)	PRESSURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	3
H406	Pressure below MIN limit (STERILIZATION phase)	PRESSURE BELOW MINIMUM LIMIT CONTACT TECH. SERVICE	3
H410	Time measurement error	INTERNAL TIMER ERROR CONTACT TECH. SERVICE	2, 3
H411	Sterilization time error	STERILIZATION TIME ERROR	3
H990	Excessive pressure (sterilization chamber, MPX)	PRESSURE BEYOND MAXIMUM LIMIT CONTACT TECH. SERVICE	2, 3
H991	Overheating (sterilization chamber, PT1)	PT1 OVERHEATING CHECK LOAD	2, 3
H992	Overheating (steam generator, PT2)	PT2 OVERHEATING CONTACT TECH. SERVICE	2, 3
H993	Overheating (layer resistance, PT3)	PT3 OVERHEATING CONTACT TECH. SERVICE	2, 3

1 = OK (warning)
2 = OK + Stopped cycle start
3 = Cycle failed + OK + RESET

24.4. SYSTEM ERRORS (CATEGORY S)

CODE	ALARM DESCRIPTION	MESSAGE ON THE DISPLAY	RESET MODE
S001	Flash memory 1 on process board failed	FLASH MEMORY NOT ACCESSIBLE CONTACT TECH. SERVICE	2, 3
S002	Flash memory 2 on process board failed	FLASH MEMORY NOT ACCESSIBLE CONTACT TECH. SERVICE	2, 3
S005	USB stick not accessible	PROBLEM WITH USB KEY CHANGE KEY	1
S006	USB stick not accessible	USB KEY NOT ACCESSIBLE CHANGE KEY	1
S007	USB stick full	USB KEY FULL CHANGE KEY	1
S009	Printer not connected	PRINTER DISCONNECTED CHECK CONNECTION	1
S010	Printer: there is no paper or there might be a configuration error	PRINTER PAPER OUT CHECK PAPER	1
S011	Printer cover open	PRINTER: DOOR OPEN	1
S012	Probable printer configuration error	PRINTER: NOT READY TRY AGAIN	1
S015	Flash memory 1 on graphical interface board	FLASH MEMORY NOT ACCESSIBLE CONTACT TECH.SERVICE	2, 3
S016	Flash memory 2 on graphical interface board	FLASH MEMORY NOT ACCESSIBLE CONTACT TECH.SERVICE	2, 3
S020	Cycle backup not done	RUN BACKUP DOWNLOAD NEW CYCLES	1
S021	Cycle storage limit exceeded	CYCLE MEMORY FULL START OVERWRITING	1
S030	Check, using a watchdog, that one of main tasks is not in crash condition	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S031	Check, using a hardware watchdog, that one peripheral is not in lock condition.	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S032	Check, using a watchdog, that one of main tasks is not in lock condition (e.g. infinite loop)	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S034	SW malfunction	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S035	SW malfunction in solenoid valve management	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S040	Check the log saving in the Flash memory	SYSTEM ERROR CONTACT TECH.SERVICE	2, 3
S041	Cycle performed with 4-minute sterilization time at 134°C	LOG SAVE ERROR CONTACT TECH. SERVICE	1
S042	Cycle performed with standard drying	4-MINUTE STERILIZATION COMPLETED	1
S050	The machine configuration has been correctly saved	CONFIGURATION SAVED REMOVE USB STICK TURN THE MACHINE OFF AND BACK ON	2, 3
S051	Process firmware and Console firmware have not been able to correctly exchange the information required for correct start	FAILED START. SWITCH OFF/ON THE DEVICE. IF THE PROBLEM PERSISTS, CONTACT THE AFTER SALES SERVICE	2, 3
S099	Error during cycle report creation	STANDARD DRYING CHECK LOAD DRYING	1
S100	SW malfunction	PROBLEM IN CREATING CYCLE REPORT CONTACT TECH. SERVICE	2, 3

1 = OK (warning)
2 = OK + Stopped cycle start
3 = Cycle failed + OK + RESET

24.5. TROUBLESHOOTING

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

24.5.1. 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	POSSIBLE CAUSE	SUGGESTED SOLUTION
E000	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 and/or the power plug pulled from the socket.	Reconnect the plug and/or turn the device on again and RESET according to the instructions.
	Network fuses blown.	Replace with good fuses of equal nominal value. (See the Summary Table in Appendix Technical Characteristics). Turn the device on again and RESET according to the instructions.
E001	Abnormal voltage peak on the mains.	Reset according to the instructions. If the problem occurs again, have the mains electric system checked by a technician.
E002	The filling tank contains water of inadequate quality.	RESET according to the instructions. Empty the filling tank and refill it with distilled water of adequate quality (<15µs/cm).
E003	The filling tank contains water of very poor quality.	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 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.*
E004	Failure to main board.	RESET according to the instructions. Contact Technical Service (see Appendix).
	Disturbance on the electrical mains.	RESET according to the instructions. If the problem occurs again, have the electrical mains checked by a technician. If the electrical mains is equipped with a Continuity system, have the system checked by a technician.
E007	One or more rear fans failed	RESET according to the instructions. Check the operation of rear fans and contact Technical Service (see Appendix).
E008	The filling/discharge tank contains water of inadequate quality.	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). 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.
E009	The filling/discharge tank contains water of very poor quality.	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 filters are present, replace the filter elements IMMEDIATELY. *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 to the device.*
E010	Door open (or not properly closed) at program start (START).	RESET according to the instructions. Properly close the door and restart the program.
	Door position microswitch failure.	Contact Technical Service (see Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
E011	The discharge tank contains water of very poor quality.	RESET according to the instructions. Empty both tanks IMMEDIATELY, 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 both tanks IMMEDIATELY, replace filter elements and fill the filling tank.
E013	The filling tank contains water of very poor quality.	RESET according to the instructions. Empty the filling tank IMMEDIATELY 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 precaution is necessary to prevent damage to the device.*
E020	Door lock mechanism limit microswitch failure. Door lock system gearmotor failure.	RESET according to the instructions. Try restarting the program a second time. If the problem persists, contact Technical Service (see the Appendix).
E021	Door lock mechanism limit microswitch failure. Door lock system gearmotor failure.	RESET according to the instructions. Contact Technical Service (see Appendix).
E022	Door lock microswitches failure	RESET according to the instructions. Contact Technical Service (see Appendix).
E030	Water level in the filling tank below minimum.	RESET according to the instructions. Top up with water up to the MAX level (or at least up to over the MIN level).
	MIN water level sensor failure.	Contact Technical Service (see Appendix).
E031	Water level in the drain tank over the MAX level.	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 Vacuum Test cycle	Run Vacuum Test as planned
E051	Reminder to run Helix Test cycle	Run Helix Test cycle as planned
E052	Reminder to run Vacuum + Helix Test combined cycle	Run Vacuum + Helix Test combined cycle as planned
E060	The autoclave cannot connect to Lan network	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).
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).
E070	Preheating activation with door open. The message is displayed after 10 minutes and after 20 minutes.	Always close the door when the sterilizer is not in cycle
E131	The selected user is not authorised to start or download the cycle	To start or download the selected cycle, ask for enabling to a user with administrator rights
E142	Console firmware version is not the correct one compared to Process firmware version. Malfunctions may occur in graphic interface management or data management to the outside (e.g. cycle reports, data on Cloud)	Update Console firmware or Process firmware to align the two firmwares to the correct version. Contact Technical Service (see Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
E900	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.
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
E901	Excessive humidity in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program.
	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.
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
E902	Excessive humidity in the sterilization chamber.	RESET according to the instructions. Thoroughly dry the inside of the chamber and restart the program.
	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 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 Appendix).
E999	Manual interruption of the sterilization or test cycle.	RESET according to the instructions.

24.5.2. 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.	
A040	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).
	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).
A101	Chamber temperature sensor failure (PT1).	Contact Technical Service (see Appendix).
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).	
	Temperature sensor short-circuit (steam generator).	
A113	Incorrect temperature sensor connection (heating element).	
	Temperature sensor short-circuit (heating element).	
A115	Temperature sensor PT5 short-circuited (conductivity measurement compensation).	
A116	ADC error.	
A117	Lack of lubrication in the door lock system	Lubricate the door lock system.
A120	Reference heating element acquisition chain fault.	Contact Technical Service (see Appendix).
A121	Reference heating element acquisition chain fault.	
A122	Reference heating element acquisition chain fault.	
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	
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	Error during Cloud firmware update	
A145	Faulty current draw detected	Check mains voltage. Contact Technical Service (see Appendix).
A201	Steam generator safety thermostat triggered.	Contact Technical Service (see Appendix)
	Steam generator or heating element malfunction.	
A202	Heating band safety thermostat triggered.	
	Heating band malfunction	
A250	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 not insert material impregnated with water or in general with liquids into the chamber.
	Drain filter obstructed.	Clean the drain filter. (See Appendix Maintenance).
	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).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
	Problem in the hydraulic circuit.	
A251	Water injection pump malfunction.	Contact Technical Service (see Appendix).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A252	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.
	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in Appendix, Technical Characteristics).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A253	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 not 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.	
A254	Water injection pump malfunction.	Contact Technical Service (see Appendix).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A255	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.
	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in Appendix, Technical Characteristics).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A256	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 not 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.	
A257	Water injection pump malfunction.	Contact Technical Service (see Appendix).
	Problem in the hydraulic circuit.	
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A258	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.
	Excessive load.	RESET according to the instructions. Check that the load does not exceed the maximum values permitted. (See the Summary Table in Appendix, Technical Characteristics).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
	Steam generator safety thermostat triggered.	
	Steam generator malfunction.	
A260	Drain filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A261	Bacteriological filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A262	Bacteriological filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
A353	Drain filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A356	Drain filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A360	Drain filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).
A362	Drain filter obstructed.	Clean the drain filter (see Appendix Maintenance).
	Problem in the hydraulic circuit.	Contact Technical Service (see Appendix).

24.5.3. HAZARDS (CATEGORY H)

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
H150	Pressure sensor failure (MPX).	Contact Technical Service (see Appendix).
H160	Pressure sensor (MPX) not properly connected to the connector.	
	Pressure sensor short-circuit (MPX).	
H400	Problem in the hydraulic circuit.	
H401	Problem in the hydraulic circuit.	
H402	Steam generator malfunction.	
	Problem in the hydraulic circuit.	
H403	Steam generator malfunction.	
	Problem in the hydraulic circuit.	
H404	Problem in the hydraulic circuit.	
	Steam generator malfunction.	
H405	Problem in the hydraulic circuit.	
	Steam generator malfunction.	
H406	Problem in the hydraulic circuit.	
	Steam generator malfunction.	
H410	Timer problem.	
H411	Sterilization time error.	
H990	General operating problem.	
H991	General operating problem.	
H992	General operating problem.	
H993	General operating problem.	

24.5.4. 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).
S005	USB stick not correctly formatted Damaged USB stick	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 stick not correctly formatted Damaged USB stick	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 stick full	Download data from USB stick or use another USB stick. 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).
S015	Error of flash memory 1 on graphical interface board. Flash memory 1 on graphical interface board failed.	Contact Technical Service (see Appendix).
S016	Error of flash memory 2 on graphical interface board. Flash memory 1 on graphical interface board failed.	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 7000 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 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 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 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 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 Appendix).
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 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).
S050	The machine configuration has been correctly saved	Turn the sterilizer off and back on.
S051	Process firmware and Console firmware have not been able to correctly exchange the information required for correct start	Contact Technical Service (see Appendix).
S099	Malfunction of control board or software	Try restarting the program a second time. Try replacing the USB key. If the problem persists, contact Technical Service (see Appendix).

CODE	POSSIBLE CAUSE	SUGGESTED SOLUTION
S100	Malfunction of control board or software	Contact Technical Service (see Appendix).

25. USER PIN RESET

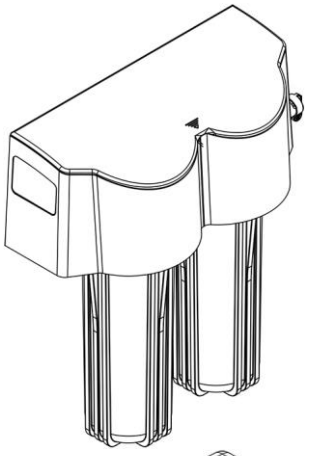
If the user enters the pin incorrectly for 3 times, it is necessary to enter the following unlock pin for four consecutive times when you will be prompted to enter pin again:

9999

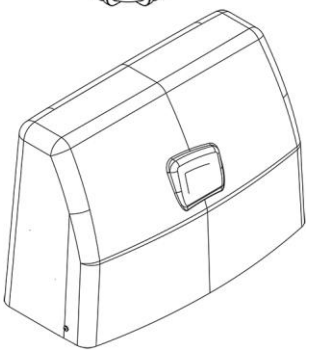
26. APPENDIX - ACCESSORIES

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

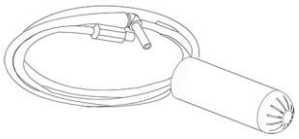
DEMINERALIZER PURE100



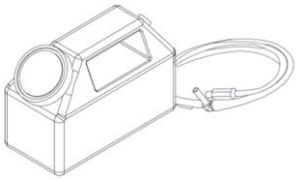
DEMINERALIZER PURE500
TWIN PURE 500



AUTOMATIC FILLING

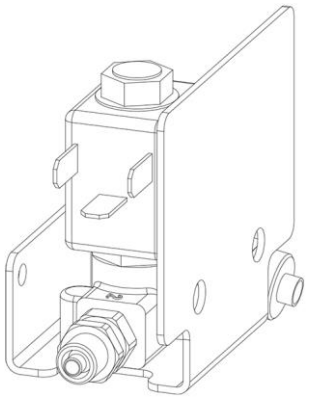


FRONT FILLING



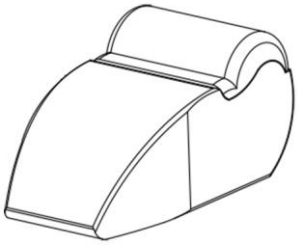
H_2O AUXILIARY SOLENOID VALVE
Additional SV kit including:

1 2-way water solenoid valve, NC - 24 V DC
2 Steel support and fastening screws
3 Connection cable with plug
4 Silicone hose with connector
5 Control valve
6 1-way valve

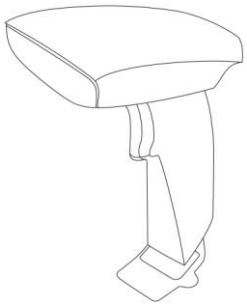


For the management of automatic filling accessories, refer to the manual of the relevant accessory.

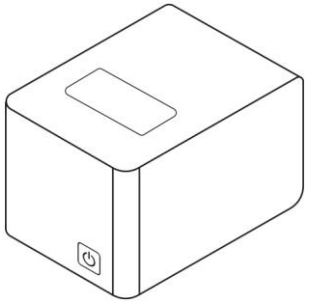
EXTERNAL PRINTER



BARCODE READER



NETWORK PRINTER

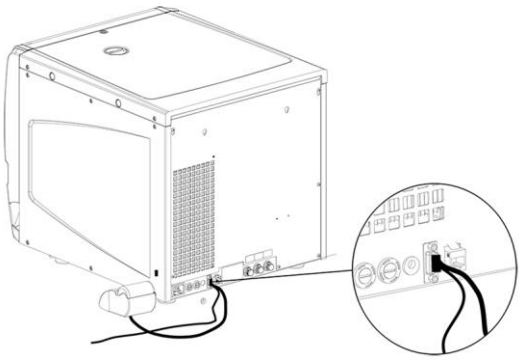


DATA STER SOFTWARE

MY TRACE SOFTWARE

27. LOCAL 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).

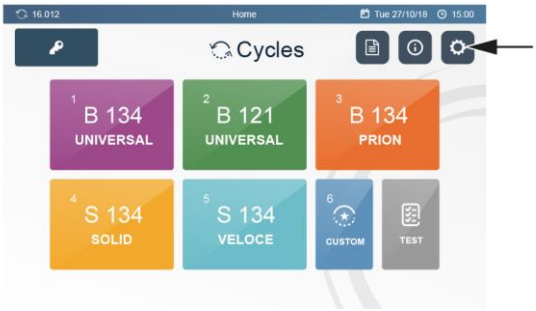
Refer to the printer manual for printer starting and paper loading.

27.1. NETWORK PRINTER CONNECTION

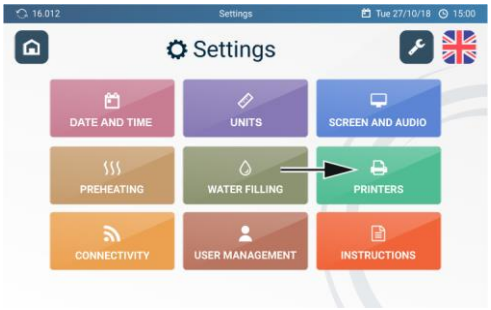
The NETWORK PRINTER allows sharing the print of labels, barecodes or reports among the various sterilizers; this function can be activated only if the NETWORK PRINTER and the sterilizers are connected to the same LAN (Local Area Network) through a Wi-Fi connection or an Ethernet cable.
For a correct installation and configuration of the NETWORK PRINTER, refer to the specific manual.
This operation is ESSENTIAL to allow the sterilizers to detect the printing device.

Once the NETWORK PRINTER installation and configuration are completed, the device can be selected on the various sterilizers through the following procedure:

Access the SETTINGS menu



Access the PRINTERS menu



Access Printer enabled to view the list of connected printers and select the NETWORK PRINTER to be enabled among them.

28. 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 l)	97400145
door gasket (28 l only)	97467176
demineralised water tank/chamber filter	97290210
discharge tank demineralisation filter and recirculation filter.	97290264

29. 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

30. 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.

www.cefla.com