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PUMA ELI R / PUMA ELI A R

EN



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1. GENERAL WARNINGS

• These instructions explain how to correctly use the following dental units:

**PUMA ELI R,
PUMA ELI A R,**

**PUMA ELI R CP,
PUMA ELI A R CP.**

PUMA ELI R CART,

PUMA ELI R ORTHO,

























PUMA ELI R CART ISO,

Please carefully read this manual before using the device.

- These instructions describe all the versions of the operating units with the maximum possible accessories, therefore not all the paragraphs are applicable to the unit you have purchased.
- It is forbidden to reproduce, store and transmit this publication by any means (electronically, mechanically, by photocopying, translating or in other ways) without a written authorisation issued by CEFLA s.c..
- The information, technical specifications and illustrations contained in this publication are not binding. CEFLA s.c. reserves the right to introduce modifications and technical improvements without having to modify these instructions.
- The manufacturer has a company policy of continual development. Although every effort is made to keep technical documentation up-to-date at all times 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 without prior notice.
- The original text is in Italian; translation from the original in Italian.
- This equipment is equipped with a device that prevents liquid back flow.

1.1. SYMBOLS

Note the meaning of the following symbols and expressions:

	Type of protection against direct and indirect contact: Class I. Type of protection against direct and indirect contact: Type B.		Equipment in compliance with essential requirements of directive 93/42/EU and subsequent changes (Class IIa Medical Equipment).
	WARNING! Failure to observe may result in equipment damage or injury to the user and/or patient.		Accessory in compliance with essential requirements of directive 93/42/EU and subsequent changes (Class I Medical Equipment).
	"Consult the instruction manual" Means that it is advisable to consult the instruction manual before using that part of the device.		Disposal symbol in accordance with Directive 2012/19/EU
	NOTE: Identifies information that is especially important for the user and/or assistant.		"Warning biological hazard". It provides information about possible risks of contamination deriving from contact with fluids, storage of infected biological waste.
	Earth ground.		Manufacturer.
	Alternating current.		Month and year of construction.
	Part sterilised in a steam autoclave up to 135° C.	SN	Device serial number.
	ON / OFF button.		DVGW mark (Quality assurance kitemark regarding supply of drinking water).
	"Refer to the instruction manual" Means that for reasons of safety you need to consult the instruction manual before using the equipment.	REF	Product/equipment identification code.
	Off (a part of the unit).		Do not push.
	On (a part of the unit).		Foot crushing hazard.
	Equipment On.		Equipment equivalent to Class 2 light source.
	Equipment OFF.		Hand crushing hazard.



1.2. INTENDED USE

- The dental units described in this manual are Medical Devices intended for dental treatment.
- The dentist's board may hold up to 5 instruments.
- The assistant's board can hold 2 suction tubes and 2 instruments.
- This equipment must be used only by adequately trained personnel (dentists and paramedics).
- The device is intended for non-continuous operation with intermittent loads (see the operating times of the individual parts in the dedicated sections).
- The device is classified as pollution degree 2.
- Overvoltage class: II.

1.2.1. CLASSIFICATION AND REFERENCE STANDARDS

• MEDICAL DEVICES classification

Classification of the dental unit in accordance with the indications given in annex IX of directive 93/42/EC as amended:

Class IIa.

• ELECTRICAL MEDICAL EQUIPMENT classification

Classification of the dental unit in accordance with standard EN 60601-1 on safety of medical equipment: **Class I - Type B.**

- Reference standards: the dental units described in this manual are designed and manufactured in compliance with IEC60601-1 3rd Ed. - 2007, IEC 60601-1-6 3rd Ed. - 2010, IEC 62366 1st Ed. - 2007, IEC 80601-2-60 1st Ed. - 2012, IEC 60601-1-2 3rd Ed., ISO 6875 3rd Ed. - 2011, ISO 7494-1 2nd Ed. - 2011 and EN 1717 (type AA and AB) standards as far as water mains safety devices are concerned.

• Classification of RADIO DEVICES AND COMMUNICATION TERMINALS (only when the WIRELESS foot control is present)

Equipment classification according to Directive 99/05/EC Art.12: **Class I.**

1.2.2. ENVIRONMENTAL CONDITIONS

The equipment is to be installed in rooms that satisfy the following requirements:

- temperature between 10 and 40 °C;
- relative humidity between 30 and 75%;
- atmospheric pressure ranging from 700 to 1060 hPa;
- altitude ≤ 3000 m;
- air pressure entering equipment ranging from 6 to 8 bar;
- water hardness at the equipment inlet must not be above 25 °f (French degrees) or 14 °d (German degrees) for untreated drinking water. For water with a higher hardness degree, it is recommended to soften water until it reaches a hardness degree between 15 and 25 °f (French degrees) or between 8.4 and 14 °d (German degrees);
- water pressure entering equipment ranging from 3 to 5 bar;
- water temperature entering equipment not higher than 25 °C.

1.2.2.1. INSTALLATION CONDITIONS

- Temperature between -10 and 70°C;
- Relative humidity between 10 and 90%;
- Atmospheric pressure: from 500 to 1060hPa.

1.2.3. WARRANTY

CEFLA s.c. stands behind its products warranting safety, reliability and performance.

The warranty is valid only under the following terms:

- Conditions given on the warranty certificate are observed.
- Yearly scheduled maintenance is performed.
- The equipment is used only as instructed in this manual.
- The electrical wiring in the room in which the equipment is installed must conform to IEC 60364-7-710 (standards for electrical wiring in medical and dental offices).
- A 3x1.5 mm² line protected by a bi-polar cut-out that conforms to applicable standards (10 A, 250 V, distance between contacts at least 3 mm) must be used to feed the equipment.



WARNING:

The colour of the three wires (POWER, NEUTRAL and EARTH) should satisfy the requirements of current standards.

- Installation, repairs, equipment enlargement, settings and, in general, any other operation requiring the casing to be opened are to be performed exclusively by personnel authorised by CASTELLINI.



1.2.4. DISPOSING THE EQUIPMENT WHEN NO LONGER USED

In compliance with Directives 2011/65/EU and 2012/19/EU regarding restriction of the use of certain hazardous substances in electrical and electronic equipment along with waste electrical and electronic equipment, it is forbidden to dispose of this equipment in the municipal waste stream as unsorted municipal waste.

When new equipment that is similar is purchased, the old equipment must be given to the dealer for disposal. As regards reuse, recycling and other forms of recovery of waste electrical and electronic equipment, the manufacturer carries out the functions defined by current local laws. A high level of separate collection of waste electrical and electronic equipment is indispensable to efficiently recycle, treat and dispose of the equipment. Recycling and treatment operations should comply with minimum standards to assure human health and high environmental protection as well as favour recycling of the materials included in the equipment. The symbol indicating separate collection for electrical and electronic equipment consists of the crossed out bin marked on the equipment.

**WARNING:**

Illegal waste clearance and disposal shall be punished as established by laws and regulations currently in force in the individual countries.

1.3. SAFETY WARNINGS

**WARNING:****• All devices are permanently installed.**

Depending on the type of dental chair the unit comes with, refer to the installation SCHEMATICS in paragraph "Technical Specifications". CEFLA s.c. shall not be held liable for any personal injury or property damage arising from failure to heed the following clause.

• Floor conditions.

The floor (continuous) should meet the load-bearing capacity set forth by DIN 1055, sheet 3.

PUMA ELI R, PUMA ELI R CP, PUMA ELI R CART, PUMA ELI R ORTHO, PUMA ELI R CART ISO dental units: the weight of the dental unit is of approx. 350 kg (including a 190 kg patient).

PUMA ELI A R, PUMA ELI A R CP dental units: the weight of the dental unit is of approx. 350 kg (including a 160 kg patient).

• This device may not be modified in any way without the authorisation of the manufacturer.

If the device is modified, appropriate examinations and tests need to be conducted in order to ensure continued safe use.

CEFLA s.c. shall not be held liable for any personal injury or property damage arising from failure to heed the following clause.

• Dental chair.

The maximum DAMA chair capacity is 190 Kg. Do not exceed this value.

The maximum DAMA A chair capacity is 160 Kg. Do not exceed this value.

• Tray holder bearing surface.

The maximum loading capacity must never be exceeded:

- tray holder attached to the dentist's board, maximum allowable load 2 Kg, evenly distributed.
- tray holder attached to the assistant's board, maximum allowable load 1 Kg, evenly distributed.
- auxiliary tray holder module, maximum allowable load on tray 3.5 Kg (without negatoscope) or 2.5 Kg (with negatoscope).

• Connections to external instruments.

The equipment can be hooked up only to other instruments that bear the CE mark.

• Electromagnetic interferences.

Use of electrical equipment that does not comply with standard IEC 60601-1 3rd Ed. - 2007 in the office or nearby may cause electromagnetic or other types of interferences resulting in dental unit malfunctions.

In these cases it is recommended to shut off the dental unit power before using this equipment.

• Replacing the drills.

Operate the turbine release and contra-angle devices only once the drill has come to a complete stop. Failure to do so, will result in damaging the locking system and drills could be released and cause injury. Exclusively use high-quality drills with a connection having a calibrated diameter. To check the conditions of the locking system, make sure the drill is firmly secured to the instrument every day before starting work. Locking system defects caused by misuse can be easily identified and are not covered by the warranty.

The drills and various instruments attached to the handpieces must comply with Biocompatibility Standard ISO 10993.

• Patients with pace makers and/or hearing aids.

When treating patients with pace makers and/or hearing aids, take into consideration the effects the instruments may have on pace makers and/or hearing aids. Carefully read technical-scientific information available on this subject.

• Implants.

If the dental unit is used for implant operations using separate equipment designed for this purpose, you are recommended to shut the power off the dental chair to avoid unwanted movements resulting from faults and/or accidental start-ups of the controls.

• Do not forget to turn off the office water supply and master switch of the equipment before leaving the surgery.**• The equipment is not protected against liquid penetration (IPX 0).****• The equipment is not suitable for use in the presence of a mixture of flammable anaesthetic gas with oxygen or nitrous oxide.****• This equipment must be stored properly so that it is kept in top working order at all times. The manufacturer shall not be held liable for misuse, carelessness or improper use of the equipment.****• The equipment may only be used by authorised and adequately trained staff (dentists and paramedics).****• The user must be present at all times when the equipment is turned on or ready for start-up. In particular, never leave the equipment unattended in the presence of children/ mentally disabled or other unauthorised personnel in general.**

Any accompanying persons must keep out of the operating area and in any case under the responsibility of the operator. The operating area refers to the space around the dental unit plus 1.5 meters.

• Quality of the water delivered by the dental unit.

The user is responsible for the quality of the water delivered by the dental unit and must adopt measures to maintain the water quality.

To ensure that delivered water is kept to quality standards, CEFLA s.c. recommends equipping the dental unit with an internal or external disinfection system.



The dental unit, once installed, is exposed to possible contaminants coming from the water mains. So, to effectively overcome this problem, it is advisable to install the dental unit only when its use will be daily and to perform the disinfecting procedures starting from the day in which it is installed by following the instructions set forth in the relevant sections.

**NOTE:**

contact your local dealer or Dentists Association for more detailed information about national laws and requirements.

Applied parts.

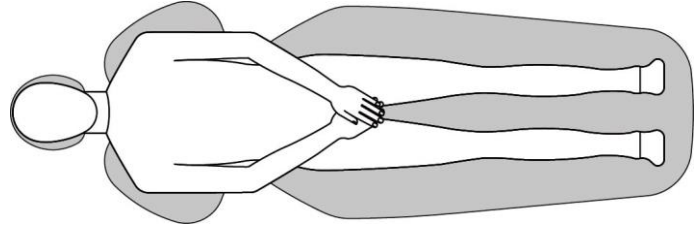
The parts of the equipment that come into contact with the patient while carrying out its functions correctly during standard use are: dental chair upholstery, armrest, curing light fibre optics, syringe terminal, disposable camera protection, scaler bits, handpiece drills, suction tube terminals.

Non applied parts that may come into contact with the patient are: dental chair armrest support, dental chair lower casing, patient-side water unit casing, water-to cup-spout, bowl, suction tubes, handpiece body.

Dental chair movement.**WARNING:**

Make sure that the patient is ready to collaborate: ask him/her to keep his/her hands and feet close to the body, avoiding incorrect postures.

Check that the patient is sitting properly when moving the dental chair (see figure).

**1.4. CLEANING AND DISINFECTION**

Cleaning is the first step of any disinfecting process.

Physically scrubbing with detergents and surface-active substances and rinsing with water removes a considerable amount of micro-organisms. If a surface is not clean first, the disinfecting process cannot be successful.

If a surface cannot be adequately cleaned, it should be protected with barriers.

The outer parts of the equipment must be cleaned and disinfected using a product for hospital use with indications for HIV, HBV and tuberculocide (medium-level disinfectant) specifically for small surfaces.

The various drugs and chemical products used in dental surgeries may damage the painted surfaces and the plastic parts. Researches and tests performed show that the surfaces cannot be fully protected against the harsh action of all products available on the market. We therefore recommend protecting with barriers whenever possible.

The harsh actions of chemical products also depend on the amount of time they are left on the surfaces.

It is therefore important not to leave the product on the surfaces longer than the time specified by the manufacturer.

It is recommended to use the specific medium-level disinfectant, **STER 1 PLUS** (CEFLA s.c.), which is compatible with:

Coated surfaces and plastic parts.**Upholstery.****WARNING:**

The **VISCOELASTIC** upholstery will stain when splashed with mordant acid. Immediately rinse with plenty of water if acid spatters on the upholstery.

Uncoated metal surfaces.

If you do not use **STER 1 PLUS**, it is recommended to use products that contain at maximum:

- **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 should be maximum 40 g per 100 g of disinfectant.

**WARNING:**

- Do not use products containing isopropyl alcohol (2-propanol, iso-propanol).
- Do not use products that contain sodium hypochlorite (bleach).
- Do not use cleaners that contain phenols.
- Do not spray the selected products directly on the surfaces.
- All products must be used as directed by the manufacturer.
- Do not mix the **STER 1 PLUS** disinfectant with other products.

**WARNING:**

The products suggested are compatible with the materials of the equipment, however damages may occur to surfaces and materials resulting from the use of different products, even if not included in the above list of excluded products.



Cleaning and disinfecting instructions.

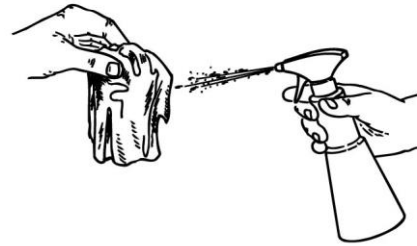
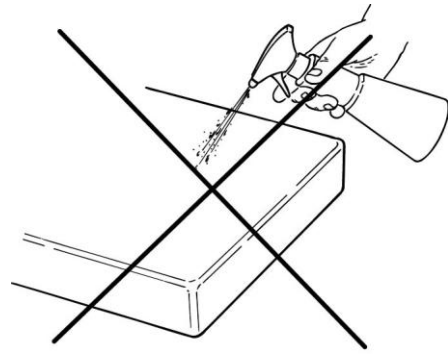
Clean and disinfect with disposable non-abrasive paper (avoid using recycled paper) or sterile gauze.

Do not use sponges or, in any case, any material that can be reused.



WARNING:

- Turn off the dental unit prior to cleaning and disinfecting the external parts.
- All materials used to clean and disinfect must be thrown away.





2. DESCRIPTION OF THE EQUIPMENT

2.1. IDENTIFICATION PLATES

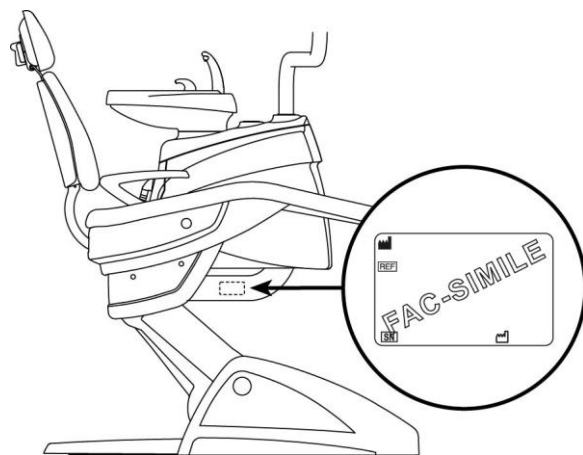
Dental unit model:

**PUMA ELI R,
PUMA ELI R CP,
PUMA ELI R CART,
PUMA ELI R ORTHO.**

The plate is found on the arm between the dental chair and water unit.

Data given on plate:

- Manufacturer's name.
- Name of equipment.
- Rated voltage.
- Type of current.
- Nominal frequency.
- Maximum absorbed power.
- Serial number.
- Month and year of manufacture.



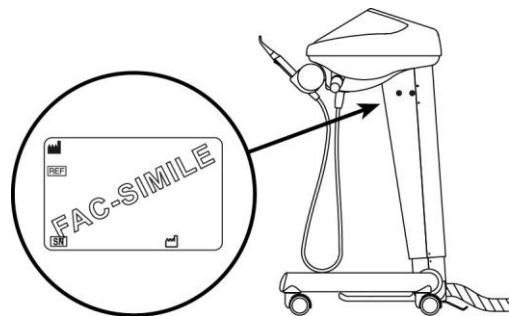
Dental unit model:

PUMA ELI R CART ISO.

The nameplate is positioned under the board.

Data given on plate:

- Manufacturer's name.
- Name of equipment.
- Rated voltage.
- Type of current.
- Nominal frequency.
- Maximum absorbed power.
- Serial number.
- Month and year of manufacture.



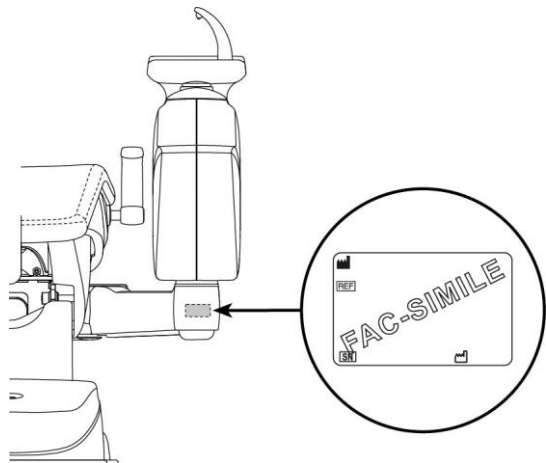
Dental unit model:

**PUMA ELI A R,
PUMA ELI A R CP.**

The plate is found on the arm between the dental chair and water unit.

Data given on plate:

- Manufacturer's name.
- Name of equipment.
- Rated voltage.
- Type of current.
- Nominal frequency.
- Maximum absorbed power.
- Serial number.
- Year of manufacture.





2.2. DENTAL UNITS

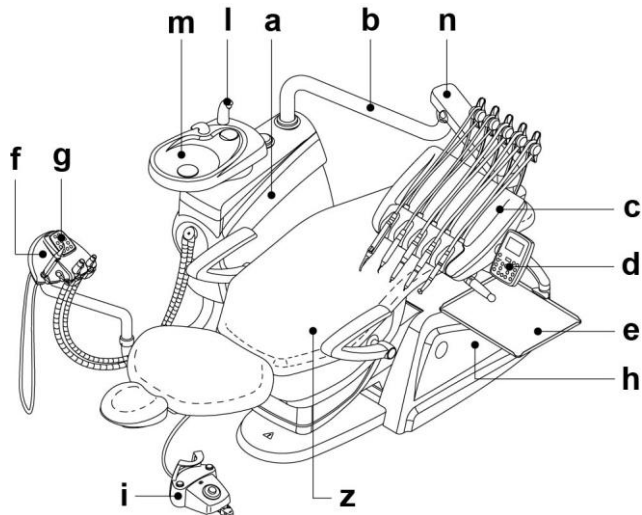
Models:

Model PUMA ELI R

RS version dentist's board (instruments will return to their original position through the pulling action of the spring-operated arms) attached to a double supporting arm, one of which is articulated and self-balancing.

Description of the different parts:

- a Water unit.
- b Adjustable arm.
- c Dentist's board.
- d Dentist's control console.
- e Tray holder.
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- n Self-balancing arm.
- z DAMA dental chair.

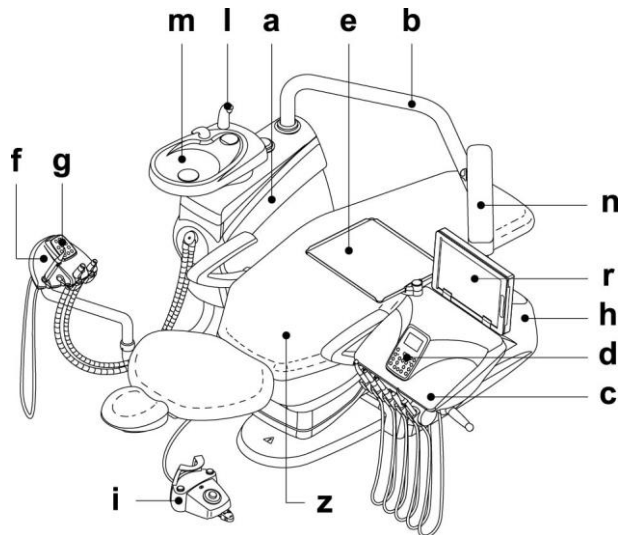


Model PUMA ELI R CP

CP version dentist's board (instruments placed vertically in housings) attached to a double supporting arm, one of which is articulated and self-balancing.

Description of the different parts:

- a Water unit.
- b Adjustable arm.
- c Dentist's board.
- d Dentist's control console.
- e Tray holder (optional).
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- n Self-balancing arm.
- r Negatoscope for panoramic X-rays (optional).
- z DAMA dental chair.

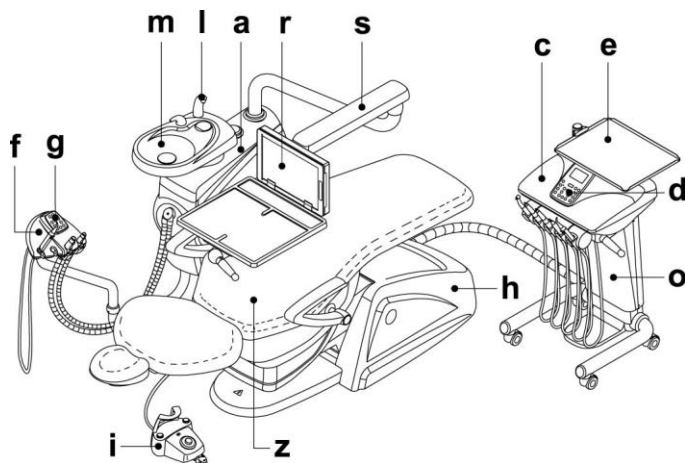


Model PUMA ELI R CART

CP version dentist's board (instruments placed vertically in housings) attached to a height-adjustable cart.

Description of the different parts:

- a Water unit.
- c Dentist's board.
- d Dentist's control console.
- e Tray holder (optional).
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- o Height-adjustable cart.
- r Negatoscope for panoramic X-rays (optional).
- s Auxiliary tray holder (optional).
- z DAMA dental chair.



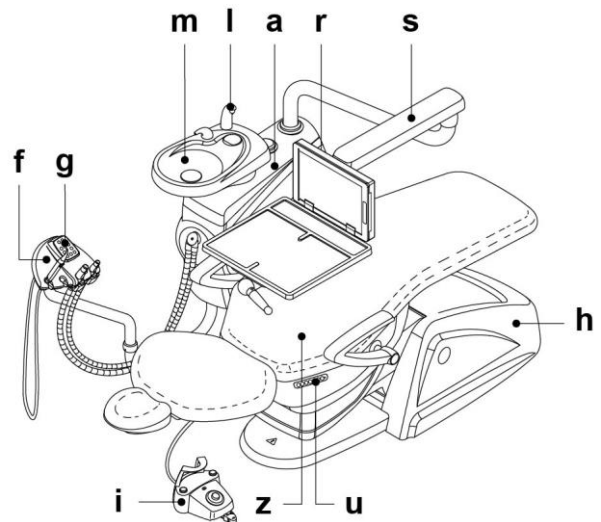


Model PUMA ELI R ORTHO

Version without dentist's board.

Description of the different parts:

- a Water unit.
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- s Auxiliary tray holder (optional).
- u Dental chair control console
- z DAMA dental chair.



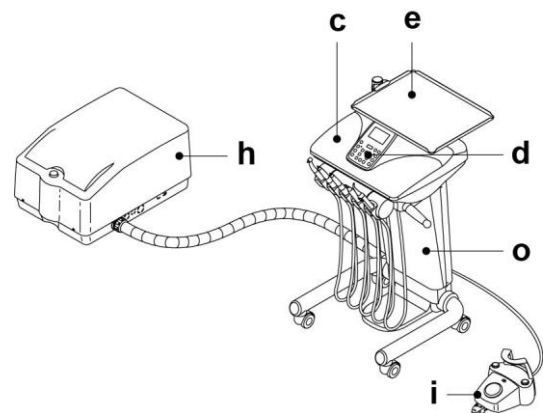
Model PUMA ELI R CART ISO

Dentist's instrument board in CP version (the instruments are vertically inserted in their seats) applied on a height-adjustable trolley and equipped with a standalone connection box.

This version is not equipped with chair and water unit.

Description of the different parts:

- c Dentist's board.
- d Dentist's control console.
- e Tray holder (optional).
- h Connection box.
- i Foot control.
- o Height-adjustable cart.



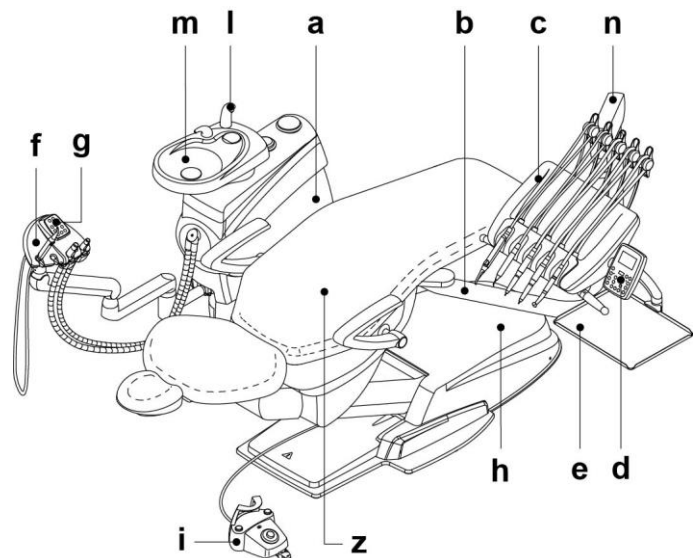
The dental units of the PUMA ELI A R series come in the following models:

Model PUMA ELI A R

RS version dentist's board (instruments will return to their original position through the pulling action of the spring-operated arms) attached to a double supporting arm, one of which is articulated and self-balancing.

Description of the different parts:

- a Ambidextrous water unit.
- b Adjustable arm.
- c Dentist's board in RS version.
- d Dentist's control console.
- e Tray holder.
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- n Self-balancing arm.
- z DAMA A dental chair.



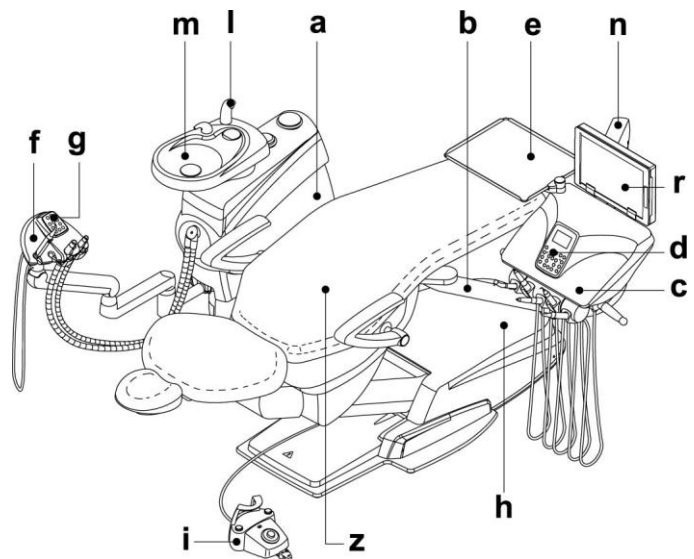


Model PUMA ELI A R CP

CP version dentist's board (instruments placed vertically in housings) attached to a double supporting arm, one of which is articulated and self-balancing.

Description of the different parts:

- a Ambidextrous water unit.
- b Adjustable arm.
- c Dentist's board in CP version.
- d Dentist's control console.
- e Tray holder (optional).
- f Assistant's board.
- g Assistant's board control console.
- h Connection box.
- i Multifunction foot control.
- l Water-to-cup spout.
- m Bowl.
- n Self-balancing arm.
- r Negatoscope for panoramic X-rays (optional).
- z DAMA A dental chair.



2.3. DENTAL CHAIR

(1) DAMA dental chair

(models PUMA ELI R - PUMA ELI R CP - PUMA ELI R CART - PUMA ELI R ORTHO)

Description of the different parts:

- a Headrest.
- b Backrest.
- c Left movable armrest (optional).
- d Right movable armrest (optional).
- e Safety foot board.

Operating time.

The operating and rest times are as follows:

work 25 sec. - rest 10 min.

Maximum permitted load.

- The maximum chair capacity is 190 Kg.
- Maximum permitted load on headrest: 30 Kg.
- Maximum permitted load on armrest: 68 Kg.



WARNING:

Do not exceed these values.

Warnings for use.

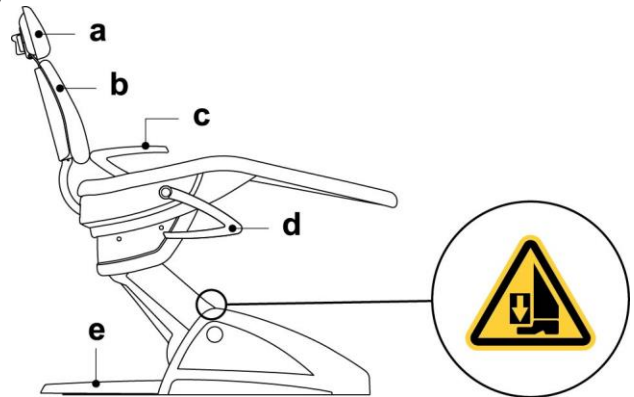


WARNING:

FOOT CRUSHING HAZARD

Pay attention to the patient and the staff during dental chair descent.

(1)



(2) DAMA A dental chair

(models PUMA ELI A R - PUMA ELI A R CP)

Description of the different parts:

- A Headrest.
- B Backrest.
- C Left movable armrest (optional).
- D Right movable armrest (optional).
- E Safety foot board.

Operating time.

The operating and rest times are as follows:

work 25 sec. - rest 10 min.

Maximum permitted load.

- The maximum chair capacity is 160 Kg.
- Maximum permitted load on headrest: 30 Kg.
- Maximum permitted load on armrest: 68 Kg.



WARNING:

Do not exceed these values.

Warnings for use.

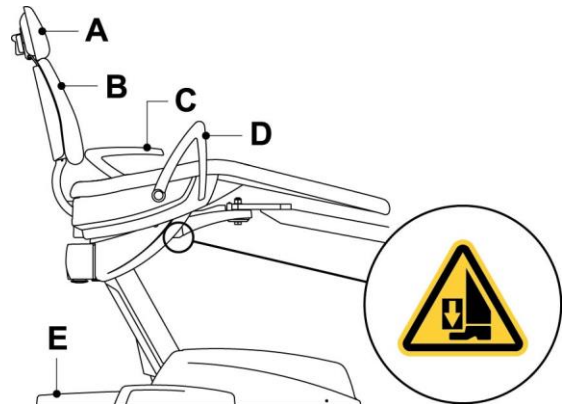


WARNING:

FOOT CRUSHING HAZARD

Pay attention to the patient and the staff during dental chair descent (on both sides).

(2)





2.4. SPECIAL WARNINGS

FOR PUMA ELI R CART and PUMA ELI R CART ISO models

During installation, the cart board can be adjusted in height to 3 predefined working positions: MIN (80 cm) - MED (90 cm) - MAX (100 cm).

NOTE:

values refer to board without tray holder.



WARNING:

Board height adjustment must be carried out by a CASTELLINI authorised technician, only.



WARNING:

When handling cart, take care to any step and/or horizontal obstacle as cart could become unstable and/or even overturn.



2.5. CONVERSION OF THE OPERATING UNIT

(Models PUMA ELI A R and PUMA ELI A R CP, only)

To convert the operating unit from the right-handed operator version to the left-handed operator version, proceed as follows:

- Move dental chair seatback to vertical position by lifting chair at least more than half its vertical travel and positioning arm and dentist's board body in central position compared to dental chair seat, so that they will not represent a side obstruction.



WARNING:

It is advisable to turn the operating unit off so as to avoid any accidental activation that could result in hazardous situations.

- Remove the cover (a) of the short arm supporting the water unit and preventing it from moving.
- Lift the element (k) locking the short arm in place.



NOTE:

slightly move arms on their pivot points so as to make element release easier.

- Turn the assistant's board arm unit until reaching a compact position with the water unit.
- Start rotating water unit and arm.



WARNING:

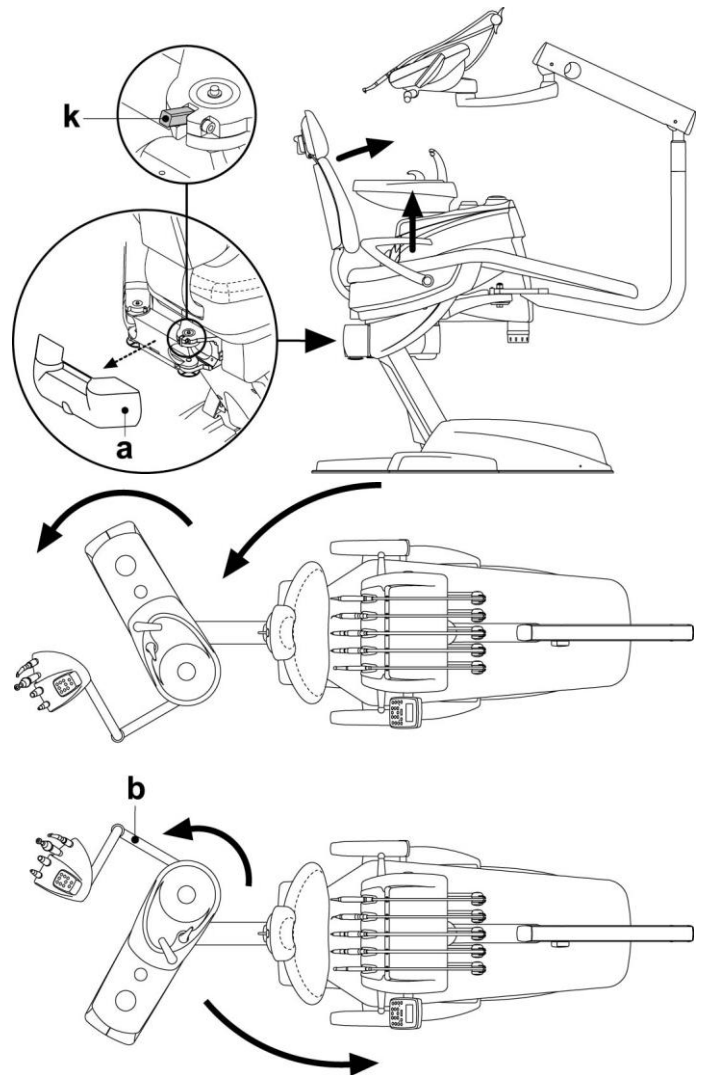
During this operation, avoid extending the arms of the operating lamp, if any, to the outside so as to prevent any excessive imbalance of the equipment.

- As soon as the water unit is aligned with the dental chair, turn the assistant's board arm unit (b) and take it to the new work position.



WARNING:

During this operation, take care not to squash instrument suction tubes and cords.





- Complete the rotation to the left of the water unit by moving it to the standard use position for left-handed operators.



NOTE:

*make sure that element (**k**) is correctly locking short arm in the new position.*

Turn bowl by hand until it reaches the new stroke end position.

- Move the pin (**c**) used to lock dentist's board arm rotation inside the opposite hole to set arm correct working area.



WARNING:

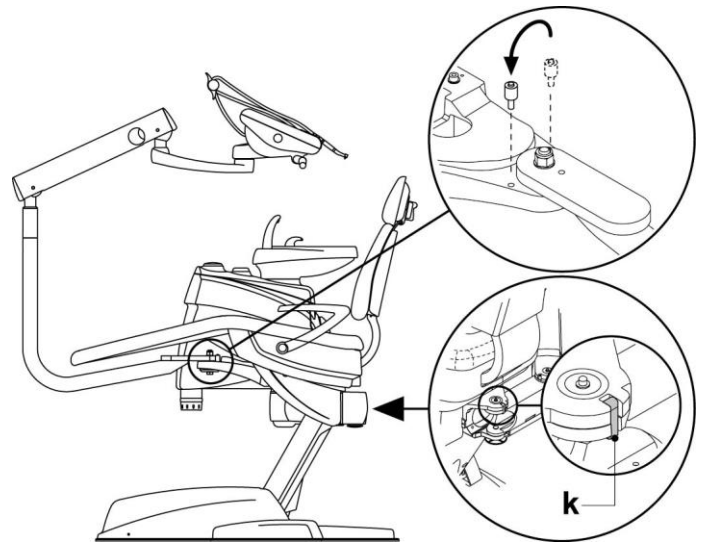
Failure to fit stroke end stop pin will make arm, during standard use, hit water unit body and thus damage it.

- Then refit cover (**a**) taking care that cover pins are duly inserted inside their seats on short arm so as to ensure water unit correct locking.
- The operating unit is now ready for use for left-handed operators.



WARNING:

Before turning the operating unit on make sure that the whole equipment is set to correct use conditions.





3. TURNING ON THE OPERATING UNIT

Dental unit model:

- PUMA ELI R
- PUMA ELI R CP
- PUMA ELI R CART

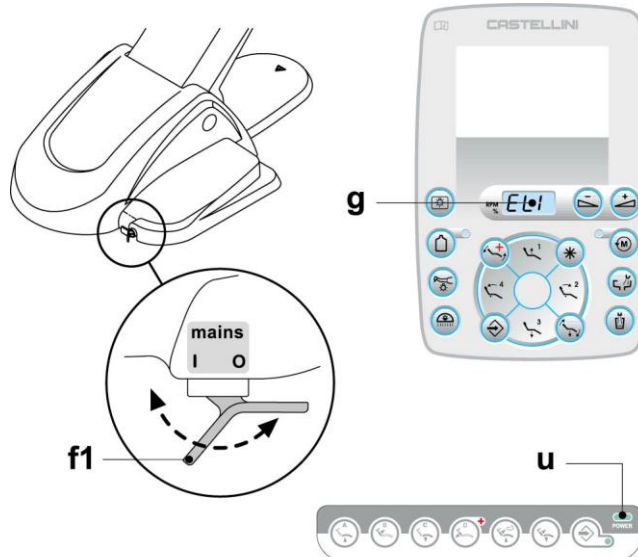
Press the main switch (f1) on the dental chair casing:

Equipment ON:

- electric system powered
- pneumatic system on
- water system on
- display (g) on.

Equipment OFF:

- electric system not powered
- pneumatic system not connected
- water system not connected
- display (g) off.



Dental unit model:

- PUMA ELI R ORTHO.

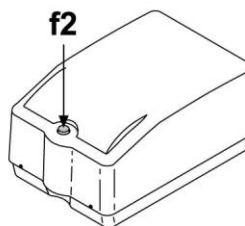
Press the main switch (f1) on the dental chair casing:

Equipment ON:

- electric system powered
- pneumatic system on
- water system on
- LED (u) On.

Equipment OFF:

- electric system not powered
- pneumatic system not connected
- water system not connected
- LED (u) Off.



Dental unit model:

- PUMA ELI R CART ISO

Press the main switch (f2) on the external connection box and check that:

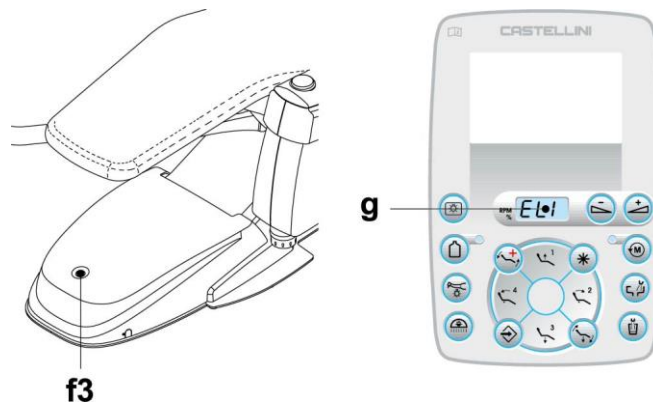
- **Display (g) is off:**
 - equipment is off
 - pneumatic system is disconnected
 - water system is disconnected.
- **Display (g) is on:**
 - equipment is on
 - pneumatic system is connected
 - water system is connected.

Dental unit model:

- PUMA ELI A R
- PUMA ELI A R CP

Press the main switch (f3) positioned on dental chair casing and check the following on the control console:

- **Display (g) is off:**
 - equipment is off
 - pneumatic system is disconnected
 - water system is disconnected.
- **Display (g) is on:**
 - equipment is on
 - pneumatic system on
 - water system is connected.



WARNING:
Main switch must be pressed with your hands.



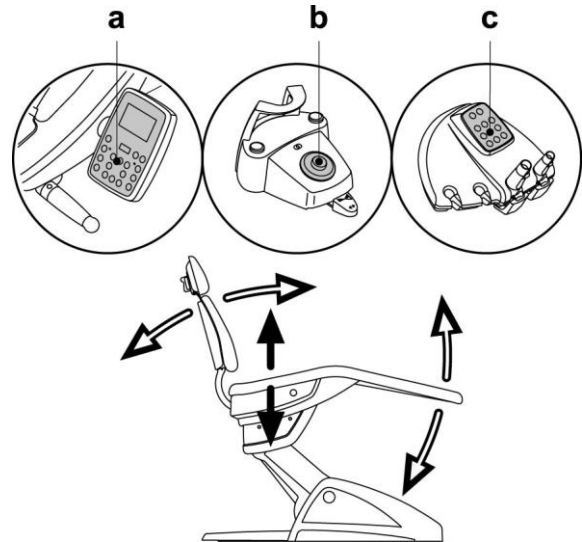
4. DENTAL CHAIR OPERATION

The dental chair can be moved as follows:

- Chair seat up/down
- Backrest rotation with inclination of the chair seat (compensated Trendelenburg).

The dental chair can be operated from the following units:

- Dentist's board (a) (see par. 5.).
- Multifunction foot control (b) (see par. 5.2.).
- Assistant's board (c) (see par. 6.).



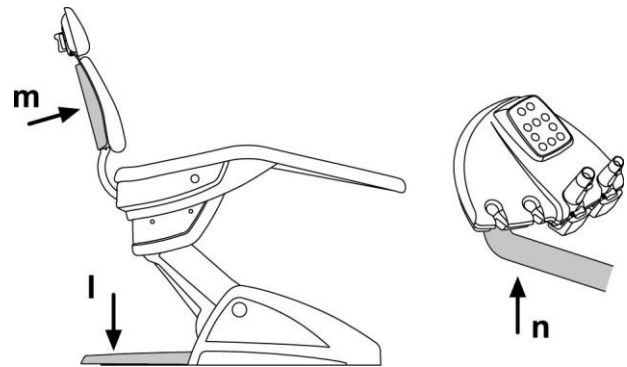
4.1. SAFETY DEVICES

The equipment is supplied with the following safety devices:

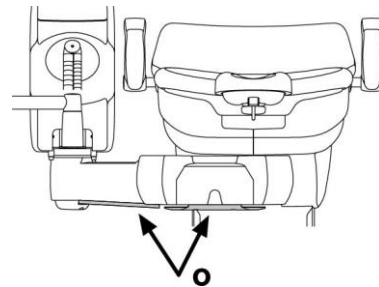
- The floor board is equipped with a device (l) that immediately stops the dental chair from moving down in the presence of an obstacle and automatically lifts it up to free the obstacle.
- The backrest is equipped with a device (m) that immediately stops the backrest from moving down in the presence of an obstacle and automatically lifts it up to free the obstacle.
- The arms of the assistant's board are equipped with a safety device (n) that immediately stops the dental chair from moving down if an obstacle is encountered and automatically moves it up to clear the obstacle.

Dental chair movements:

- with the instrument extracted NOT working: manual movements allowed, automatic movements inhibited, but if they are already in progress at the moment of extraction they are not interrupted;
- with instrument extracted and working: all the chair movements are inhibited.



- Dental units model PUMA ELI A R and PUMA ELI A R CP: water unit supporting arms are equipped with a safety device (o) that immediately stops the dental chair from moving down if an obstacle is encountered and automatically moves it up to clear the obstacle.





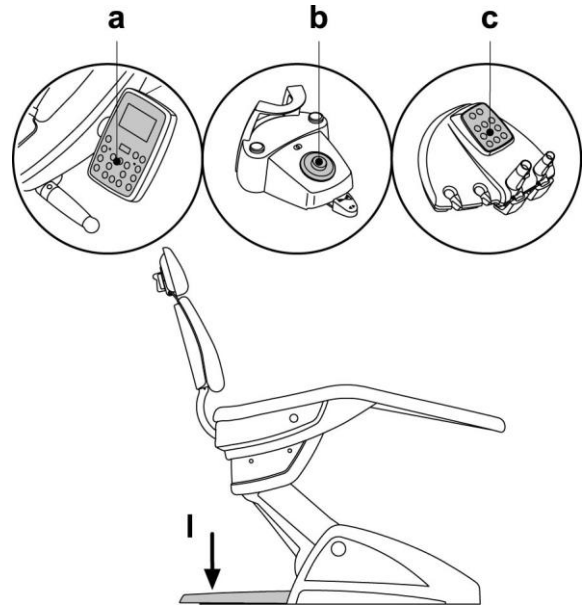
4.2. EMERGENCY DEVICES



WARNING:

Use the devices below when movement of the equipment needs to be stopped:

- **Dental chair control buttons (a) or (c).**
By pressing any of the dental chair control buttons all movements will be stopped.
- **Foot control (b).**
When the foot control is activated, all movements of the equipment are stopped.
- **Foot board (i).**
When the chair foot board is actuated, all movements of the equipment are blocked.



4.3. ADJUSTABLE HEADREST

The headrest may be of two types:

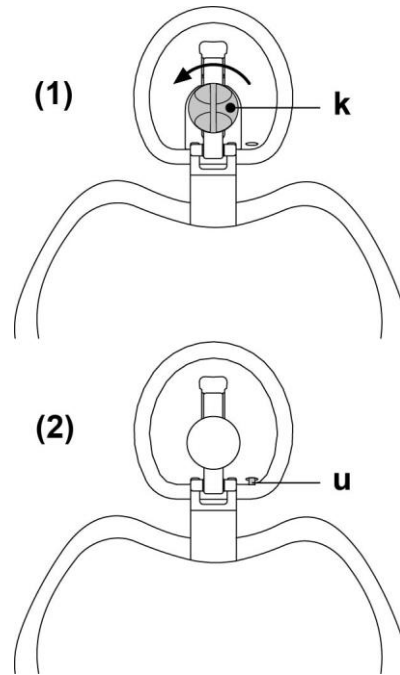
- (1) with manual cushion lock lever
- (2) with pneumatic cushion lock lever

Adjusting headrest height.

- with manual lock (1):
The head rest blade is positioned through a magnetic clutch. The operator should pull up and/or push down the headrest until it is in the desired position.
- with pneumatic lock (2):
Press and hold down the locking button (u) to position the headrest as desired. Once you have reached the desired position, release the button (u) to lock the headrest in place.

Adjusting the cushion.

- with manual lock (1):
Rotate the lock knob (k) anti-clockwise, position the cushion as desired and then re-tighten the lock knob.
- with pneumatic lock (2):
Press the lock button (u) and keep it pressed as you adjust the cushion as desired. Once the cushion is oriented as desired just release the button (u) to lock in place.



Proper positioning of the headrest.

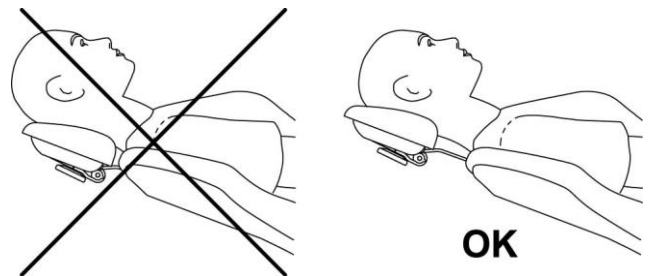


WARNING:
For correct use of the headrest, position the patient's head as shown in the figure.

Warnings for use.



- WARNING:**
- Maximum permitted load on headrest: 30 Kg.
 - Do not attempt to move headrest while patient is resting against it.
 - Do not attempt to modify the position of the cushion without first releasing the lock mechanism.
 - The pneumatic locking device is active only when the air circuit is pressurized and the dental unit is on.





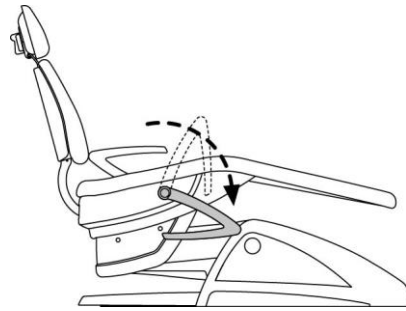
4.4. MOVABLE ARMRESTS (OPTIONAL)

Both armrests can be turned clockwise downwards so that the patient can more conveniently get on and off the chair.

Safety warnings.











WARNING:
Maximum load supported by armrest: 68 kg.




4.5. DENTAL CHAIR CONTROL PAD (MODEL PUMA ELI R ORTHO, ONLY)


Description of the buttons:

-  Dental chair function save button (with relevant activation LED).
-  Automatic return position recall button.
-  Rinsing position recall button.
-  Seat up and programmed position "A" recall button.
-  Backrest up and programmed position "B" recall button.
-  Seat down and programmed position "C" recall button.
-  Backrest down and programmed position "D" recall button.
-  Equipment On LED.




-  **NOTE:**
operation of dental chair buttons:
 - Button pressed shortly: automatic recall of set position.
 - Button held pressed: manual positioning.

Setting the dental chair "Rinse Position" and "Automatic Return".
See paragraph 5.1.1.

-  **NOTE:**
storage mode activation is signalled by a short sound (BEEP) and by the turning on of SAVE button LED.

Programming the chair positions A, B, C and D.
See paragraph 5.1.2.

-  **NOTE:**
storage mode activation is signalled by a short sound (BEEP) and by the turning on of SAVE button LED.



5. DENTIST'S BOARD OPERATION

Layout of instruments.

The positions of the instruments on the board are determined by the customer at the time of order.

Activation of instruments.

- The syringe is always on (see paragraph 5.3.).
- The curing light is turned on with the relevant key, when the instrument is withdrawn (see paragraph 5.7.).
- Intraoral camera turns on when the instrument is extracted (see paragraph 5.8.).
- Once extracted, all other instruments are operated with the foot control (see paragraph 5.2.).

Simultaneous use of the instruments.

An interlocking device ensures that the instruments are not used simultaneously.


The first extracted instrument is ready to be used while those extracted later are disabled by the interlocking device.

This interlocking device allows the drill to be changed in one instrument while another is used on the patient.

Positioning the dentist's board.

The dentist's board can move in all directions.

To adjust the height of the board and/or direct it horizontally, simply grasp the handle (**a**).

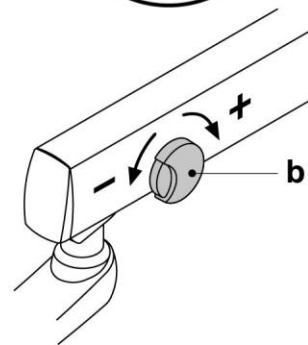
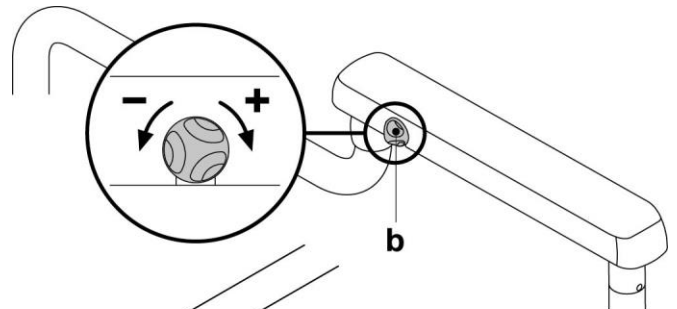
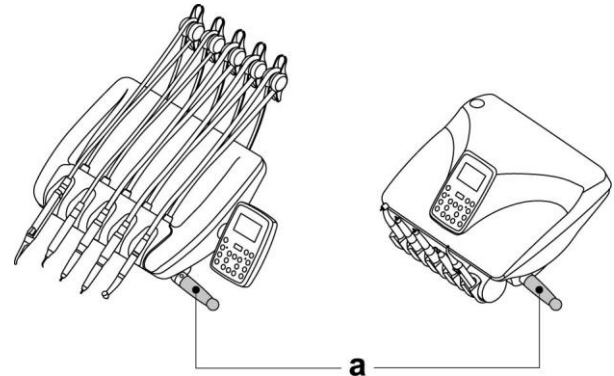
 **NOTE for models PUMA ELI R CART:**
dentist's board height can be set only during equipment installation (see paragraph 2.4).

Pantograph arm adjustment.

Balancing of the pantograph arm is determined at the time of installation. Any future adjustments can be made using the knob (**b**) on the pantograph arm.

Clockwise rotation: to increase clutch resistance of the pantograph arm.

Counter-clockwise rotation: to decrease clutch resistance of the pantograph arm.

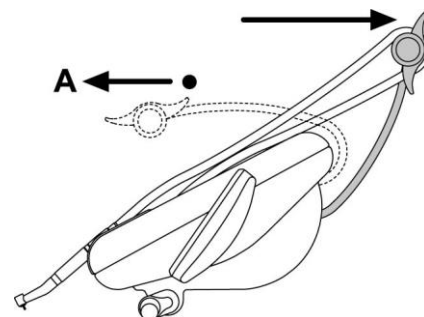


Instrument return arm lock device (only for RS version).

If this device is provided, the instrument recall arm can be locked in the instrument extracted position.

Lock engages with a click at about 2/3 of the total arm travel.

To go back to the original condition (unlock), simply move the arm to the end of its travel (**A**).



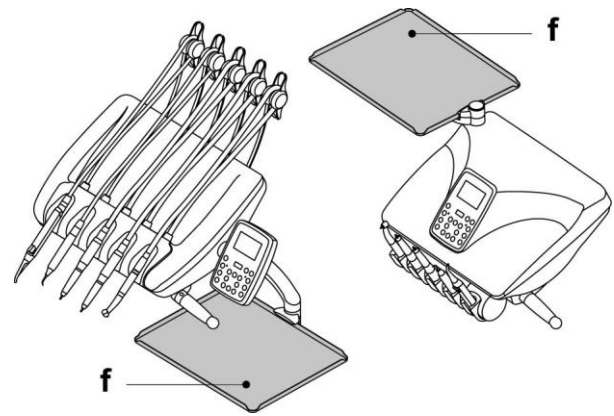


Tray holder.

The tray holder module (f) is made of stainless steel and can easily be removed from its support.



WARNING:
Maximum permitted load on the tray holder (f): 2 kg evenly distributed.



Cleaning the dentist's board handle.

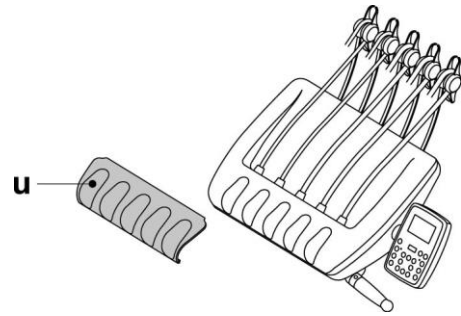
Clean the dentist's module handle using a suitable product (see Paragraph 1.4).

Dentist's board cleaning.

Clean the dentist's board using a suitable product (see Paragraph 1.4).



NOTE for RS version instrument boards:
the silicone instrument holder (u) can be sterilized in an autoclave at 121°C (rubber cycle).



Instrument cords.

Clean the instrument cord using a suitable product (see Paragraph 1.4).





















WARNING:
To remove cord, contact the Technical Service.



5.1. DENTIST'S CONTROL CONSOLE


Description of the buttons:

-  INCREASE button: increases the settable values.
-  DECREASE button: decreases the settable values.
-  Fibre optics lighting ON/OFF button.
-  Micromotor rotation direction inversion control button with relevant operating LED.
-  Operating light on/off button.
-  Water-to-cup button.
-  Water-to-bowl button.
-  S.S.S. system ON/OFF button with relevant operating LED.
-  Negatoscope on/off button on console.
-  Dental chair function save button.
-  Automatic return position recall button.
-  Rinsing position recall button.
-  Emergency position recall button.
-  Seat up and programmed position 1 recall button.
-  Backrest up and programmed position 2 recall button.
-  Seat down and programmed position 3 recall button.
-  Backrest down and programmed position 4 recall button.

 **NOTE: operation of dental chair buttons.**

- Button pressed shortly: automatic recall of set position.
- Button held pressed: manual positioning.

Indicator lights:

 Display for viewing:

- power or speed of the dynamic instruments,
- storage function enabled,
- AUTOSTERIL-ST or TIME FLUSHING cycle activation.





5.1.1. SETTING THE DENTAL CHAIR "AUTOMATIC RETURN" AND "RINSE POSITION"

- Bring the dental chair into the desired position with the manual movement buttons.
- Hold button "SAVE" for at least 2 seconds to activate save mode. Storage mode activation is signalled by a short beep and by the flashing 0 on the console display.



NOTE:

hold down button SAVE for at least 2 seconds to quit without saving the changes made.

- Press the "Automatic Return" or "Rinsing Position" buttons to associate the position with the button (e.g. "Rinsing Position").



NOTE:

the number for the selected button lights up on the display to confirm the position has been saved:

5 = AUTOMATIC RETURN

6 = RINSE POSITION

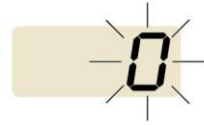


NOTE:

the "Rinsing Position" button brings the chair backrest into the rinse position without changing the position of the chair seat. When the button is pressed again, the backrest goes back to the previous position.



>2 sec



5.1.2. DENTAL CHAIR POSITION PROGRAMMING

- Bring the dental chair into the desired position with the manual movement buttons.
- Hold button "SAVE" for at least 2 seconds to activate save mode. Storage mode activation is signalled by a short beep and by the flashing 0 on the console display.



NOTE:

hold down button "SAVE" for at least 2 seconds to quit without saving the changes made.

- Press the 1, 2, 3 or 4 buttons to associate the relevant position to the button (e.g. 3).

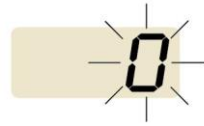


NOTE:

the number of the button selected (e.g. 3) will be shown on the console display to confirm that it has been stored.



>2 sec



5.1.3. EMERGENCY BUTTON

This button can be used in the event of an emergency to bring the patient into the Trendelenburg position.



NOTE:

the Trendelenburg position is already set and cannot be changed.



5.1.4. TURNING ON THE NEGATOSCOPE

Press the ON/OFF button.

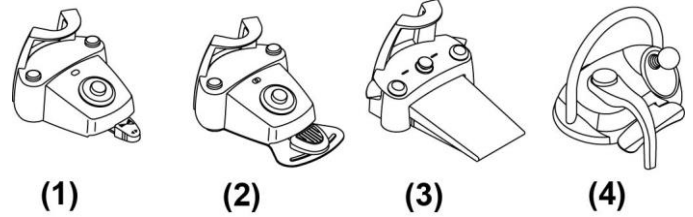




5.2. FOOT CONTROL

4 types of foot controls are available:

- (1) "Multifunction" foot control (see paragraph 5.2.1).
- (2) "Pressure operated" foot control (see paragraph 5.2.2).
- (3) "Power pedal" foot control (see paragraph 5.2.3).
- (4) "Lateral excursion" foot control (see paragraph 5.2.4).



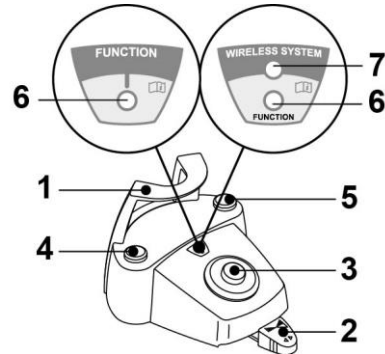
NOTE:

the "side travel" and "push-pedal" foot controls can also be supplied in WIRELESS version (see paragraph 5.2.5).

5.2.1. "MULTIFUNCTION" FOOT CONTROL

Description of the parts.

- 1 Handle.
- 2 Control pedal.
- 3 Dental chair movements.
- 4 Chip-air/Patient rinsing position recall control.
- 5 Water Clean System/Automatic dental chair return control.
- 6 LED (not active).
- 7 Battery charge LED (wireless version only).



Control pedal (2).

With instrument removed

- Starts the instrument.
- Adjusts the rpm of rotary instruments.
- To the right: operation with spray (if foreseen for selected instrument).



NOTE:

at the end of work, air is automatically blown to eliminate any drops of liquid remaining in the spray ducts.



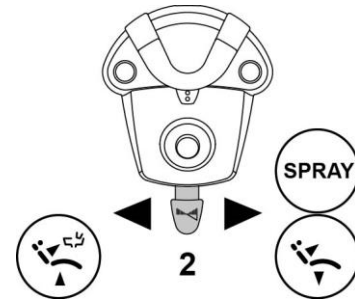
WARNING:

These dental chair functions are activated by keeping the pedal at the end of the travel for at least 2 seconds.



NOTE:

these functions are NOT active in PUMA ELI R ORTHO models.



Dental chair movement Joystick operation (3).

It controls the following movements:

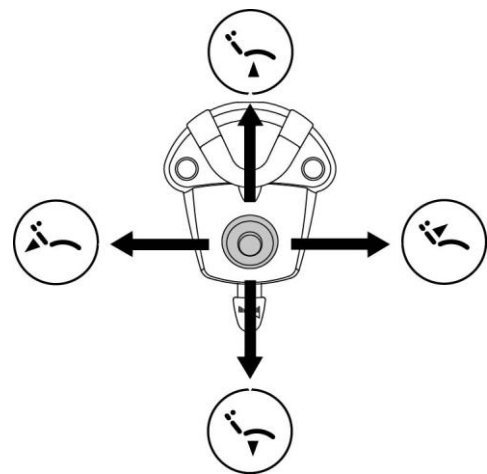
- Dental chair seat up.
- Dental chair backrest up.
- Dental chair seat down.
- Dental chair backrest down.

To stop movement, release the control.



NOTE:

all the buttons used to move the dental chair are inoperative when an instrument is removed and the foot control pedal is actuated.





Left-hand button operation (4).

- Holding down the button for at least 2 seconds with instrument extracted:
Chip-air control: sends a jet of air to the Turbine or the Micromotor. Air flow is activated by pressing the button; the jet of air is interrupted as soon as button is released.
- Holding down the button for at least 2 seconds with the instruments in rest position:
activation of the "Patient rinsing position" program.



NOTE:

pressing the button a second time returns dental chair back into working position.



NOTE:

this function is NOT active in PUMA ELI R ORTHO models.

Right-hand button operation (5).

- Holding down the button for at least 2 seconds with instrument extracted:
Water Clean System Control: sends a jet of running water to instruments such as the Turbine, the Micromotor and the Scaler for rinsing the spray ducts. Water is delivered by pressing the button. Water is no longer delivered as soon as the button is released and air is automatically blown to eliminate any drops of liquid remaining inside the spray ducts.
- Holding down the button for at least 2 seconds with the instruments in rest position:
Activation of the "Automatic dental chair return" programme.

Wireless version.

This foot control can also be supplied in wireless version (see Paragraph 5.2.4).

Protection against liquid penetration.

The foot control is protected against liquid penetration. Degree of protection: IPX1.

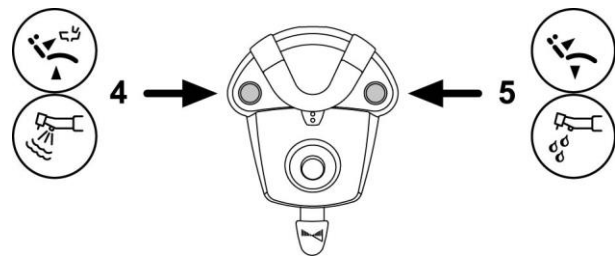
Cleaning.

Clean the foot control using a suitable product (see Paragraph 1.4).



NOTE:

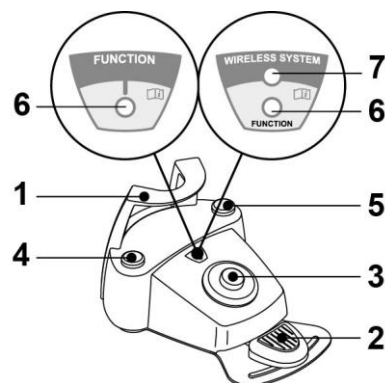
if the foot control slips on the floor, remove any dust from the slip-proof rubber found under the base with a dry cloth.



5.2.2. "PUSH-PEDAL" FOOT CONTROL

Description of the parts.

- 1 Handle.
- 2 Control pedal.
- 3 Dental chair movements.
- 4 Chip-air/Patient rinsing position recall control.
- 5 Water Clean System/Automatic dental chair return control.
- 6 Spray operation LED.
- 7 Battery charge LED (wireless version only).





Control pedal (2).

Operation:

- Remove the instrument.
- Push the control pedal (a) to start the instrument.
- Adjust the rpm/power of the instrument with the control pedal:
 - to the right: increase;
 - to the left: decrease.



NOTE:

the control pedal adjusts the speed/power of the instrument from the minimum to the maximum value set from the dentist's board.

- To stop the instrument, simply release the control pedal (a).



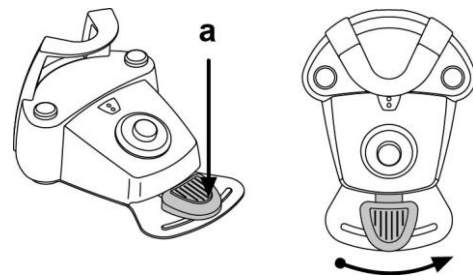
NOTE:

with the spray active, at the end of the operation a blast of air is automatically activated to remove any residual drops of liquid in the spray ducts.



WARNING:

Instrument spray is activated and deactivated by pressing the buttons (4) or (5).
A short acoustic signal warns of the switching.
When the LED (6) is on, it indicates operation with spray.



Dental chair movement Joystick operation (3).

It controls the following movements:



Dental chair seat up.



Dental chair backrest up.



Dental chair seat down.



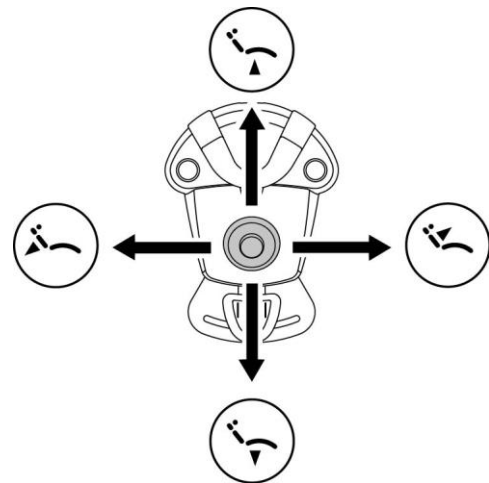
Dental chair backrest down.

To stop movement, release the control.



NOTE:

all the buttons used to move the dental chair are inoperative when an instrument is removed and the foot control pedal is actuated.



Left-hand button operation (4)

Operation:

- Holding down the button for at least 2 seconds with the instruments in rest position:

Activation of the "Patient rinsing position" program.



NOTE:

pressing the button a second time returns dental chair back into working position.



NOTE:

these functions are NOT active in PUMA ELI R ORTHO models.

- Holding down the button for at least 2 seconds with instrument extracted:

Chip-air control: sends a jet of air to the Turbine or the Micromotor.

Air flow is activated by pressing the button; the jet of air is interrupted as soon as button is released.



NOTE:

the control works only when the Turbine and Micromotor are in working position.

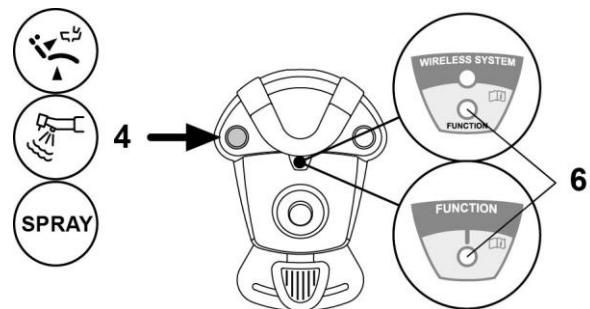
- Press the button shortly with the instrument extracted:

Activation or deactivation of instrument spray.



WARNING:

A short acoustic signal warns of the switching.
When the LED (6) is on, it indicates operation with spray.





Left-hand button operation (5).

Operation:

- Holding down the button for at least 2 seconds with the instruments in rest position:

Activation of the "Automatic dental chair return" programme.



NOTE:

these functions are NOT active in PUMA ELI R ORTHO models.

- Holding down the button for at least 2 seconds with instrument extracted:
Water Clean System Control: sends a jet of running water to instruments such as the Turbine, the Micromotor and the Scaler for rinsing the spray ducts.

Water is delivered by pressing the button. Water is no longer delivered as soon as the button is released and air is automatically blown to eliminate any drops of liquid remaining inside the spray ducts.

- Press the button shortly with the instrument extracted:

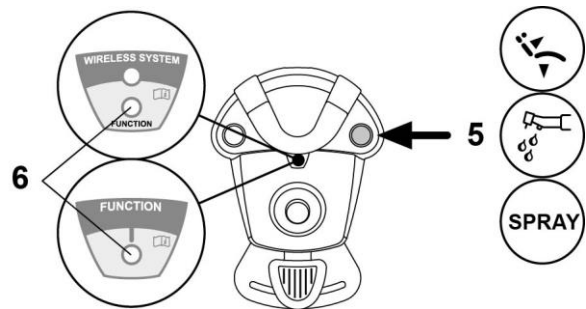
Activation or deactivation of instrument spray.



WARNING:

A short acoustic signal warns of the switching.

When the LED (6) is on, it indicates operation with spray.



Wireless version.

This foot control can also be supplied in WIRELESS version (see "WIRELESS foot control" paragraph).

Protection against liquid penetration.

The foot control is protected against liquid penetration. Degree of protection: IPX1.

Cleaning.

Clean the foot control using a suitable product (see Paragraph 1.4).



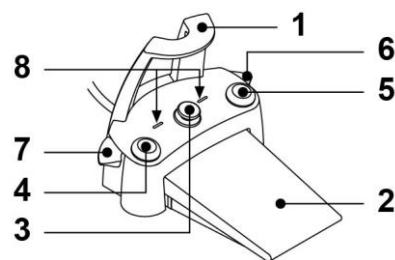
NOTE:

if the foot control slips on the floor, remove any dust from the slip-proof rubber found under the base with a dry cloth.

5.2.3. "POWER PEDAL" FOOT CONTROL

Description of the parts.

- 1 Handle.
- 2 Foot control.
- 3 Dental chair movements.
- 4 Chip-air control or activation/deactivation of instrument spray function.
- 5 Water Clean System control or activation/deactivation of instrument spray function.
- 6 Programme "B" recall or automatic dental chair return activation.
- 7 Programme "A" recall or patient rinse position activation.
- 8 Spray operation LED.



Foot control operation (2).

- With instrument removed

- Pushing the pedal (a), the instrument is started.
The instrument rpm (or power) can be adjusted by varying the pressure exerted on the foot control.



NOTE:

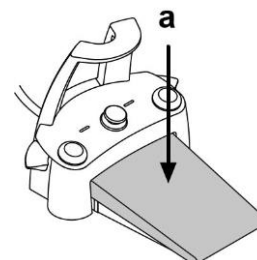
the foot control adjusts the speed/power of the instrument from the minimum to maximum value set from the instrument board.

- Release the foot control to stop instrument operation.



NOTE:

with the spray active, at the end of the operation a blast of air is automatically activated to remove any residual drops of liquid in the spray ducts.







- With instrument in rest position

When the foot control is pressed, all automatic dental chair movements are automatically blocked.



Dental chair movement Joystick operation (3).

It controls the following movements:

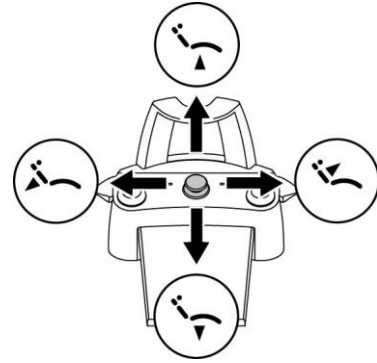
-  Dental chair seat up.
-  Dental chair backrest up.
-  Dental chair seat down.
-  Dental chair backrest down.

To stop movement, release the control.



NOTE:

all dental chair movements are blocked when an instrument is being used or the BIOSTER system is running.



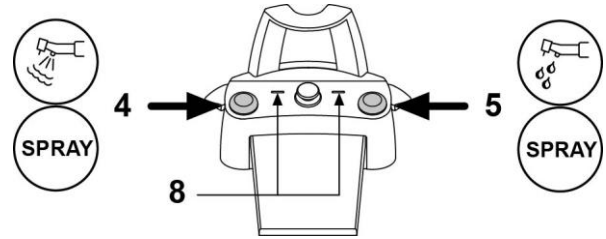
Left-hand button operation (4).

- Holding down the button for at least 2 seconds with instrument extracted:
Chip-air control: sends a jet of air to the Turbine or the Micromotor. Air flow is activated by pressing the button; the jet of air is interrupted as soon as button is released.
- Press the button shortly with the instrument extracted:
Activation or deactivation of instrument spray.



WARNING:

**A short acoustic signal warns of the switching.
When the LED (8) is on, it indicates operation with spray.**



Right-hand button operation (5).

- Holding down the button for at least 2 seconds with instrument extracted:
Water Clean System Control: sends a jet of running water to instruments such as the Turbine, the Micromotor and the Scaler for rinsing the spray ducts. Water delivery is activated by pressing the button (5); when the button is released, the jet of water is interrupted and a blast of air is automatically activated to remove any residual drops of liquid in the spray ducts.
- Press the button shortly with the instrument extracted:
Activation or deactivation of instrument spray.



WARNING:

**A short acoustic signal warns of the switching.
When the LED (8) is on, it indicates operation with spray.**

Right-hand lever operation (6).

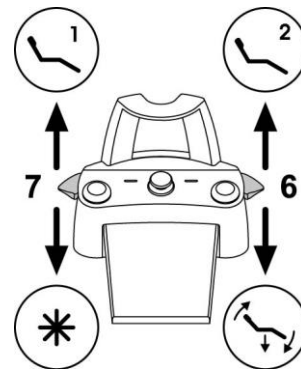


NOTE:

the lever functions only if the instruments are in rest position. This function is NOT active in the PUMA ELI R ORTHO models.

For safety reasons, the selected function starts only after the lever has been briefly actuated and then released.

- Lever pushed downwards:
"Automatic dental chair return" programme activated.
- Lever pulled upwards:
Dental chair program "2" start.



Left-hand lever operation (7).



NOTE:

the lever functions only if the instruments are in rest position. This function is NOT active in the PUMA ELI R ORTHO models.

For safety reasons, the selected function starts only after the lever has been briefly actuated and then released.

- Lever pushed downwards:
Activation of the "Patient rinsing position" programme.



NOTE:

when the lever is actuated the second time, the dental chair reaches its work position.

- Lever pulled upwards:
Dental chair program "1" start.



Protection against liquid penetration.

The foot control is protected against liquid penetration. Degree of protection: IPX1.

Cleaning.

Clean the foot control using a suitable product (see Paragraph 1.4).



NOTE:

if the foot control slips on the floor, remove any dust from the slip-proof rubber found under the base with a moist cloth.

5.2.4. "SIDE TRAVEL" FOOT CONTROL

Description of the parts.

- 1 Handle.
- 2 Horizontal movement control lever.
- 3 Joystick for dental chair movements.
- 4 Vertical movement control.

Control pedal (2).

With instruments in rest position, control lever (2) has 2 different operating modes, based on whether the vertical control (4) is pressed or not.



NOTE:

once control (4) is pressed, 5 seconds are allowed to operate lever (2); after this time, control will be automatically reset.

Operation with instruments in rest position

- Lever (4) in end-of-travel position to the right: operating lamp turning on/off.

Operation with instruments in rest position after having pressed control (4)

- Lever (2) in end-of-travel position to the right: dental chair automatic return.
- Lever (2) in end-of-travel to the left: patient rinsing position recall. If the lever (2) is pushed all the way to the left, the dental chair reaches the work position.



NOTE:

these dental chair functions are activated by keeping the pedal at the end of the travel for at least 2 seconds.



NOTE:

these functions are NOT active in PUMA ELI R ORTHO models.

Operation with instrument removed

- Operating lever (2) to the right: instrument is started and its speed/position is adjusted.



NOTE:

if the removed instrument is a camera, the freeze function is carried out.

- Lever (2) in end-of-travel to the left: the CHIP AIR is activated for Turbine or Micromotor. The air jet is interrupted as soon as lever (2) is released.



NOTE:

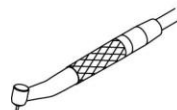
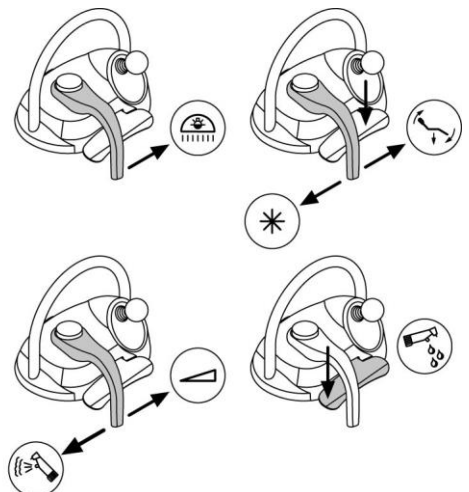
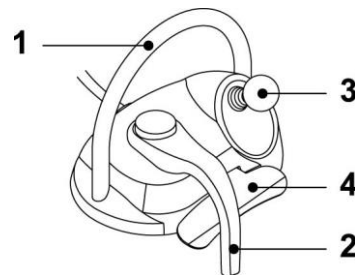
if the camera is withdrawn, the saved images can be selected one after the other.

- If control (4) is held down, CHIP WATER will be activated for turbine, micromotor or scaler. The water jet is interrupted as soon as control (4) is released.



NOTE:

if the removed instrument is a camera, the full-screen displaying of the selected image will be activated/deactivated.





Activation/deactivation of spray water to instruments.

Spray is delivered by keeping control (4) pressed with the instrument active (lever (2) operated)

NOTE:

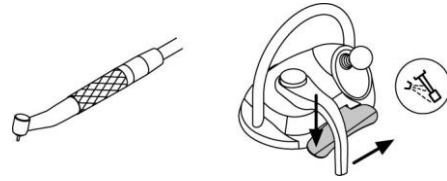
the foot control can be set so that when the instrument is selected and lever (2) is in rest position, pressing the button (4) will enable operation to always be with spray (ON/OFF). This configuration change of the foot control can be carried out only by a CASTELLINI Authorised Technician.

NOTE:

the type of delivered spray can be set through the dentist's console.

NOTE:

at the end of work a blast of air is automatically activated to eliminate any residual drops of liquid in the instrument ducts.



Joystick for dental chair movement (3).

Joystick for dental chair movement (3) has 2 different operating modes, based on whether the vertical control (4) is pressed or not.

NOTE:

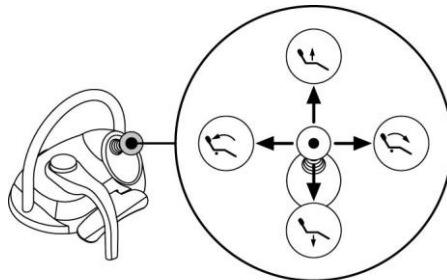
once the pedal (4) is pushed, the joystick (3) must be actuated within 5 seconds. The function is automatically cancelled once this time has elapsed.

Operation with instruments in rest position

Joystick (3) controls the 4 manual movements of the dental chair:

NOTE:

release the joystick to stop manual movement.

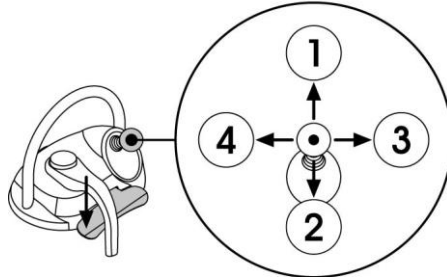


Operation with instruments in rest position after having pressed control (4)

Joystick (3) recalls the 4 programmes of the dental chair.

NOTE:

to interrupt automatic movement, simply operate joystick again.



With instrument removed but not active

Joystick (3) controls the 4 manual movements of the dental chair.

NOTE:

under these conditions, the 4 programmes are blocked.

With instrument active (lever (2) operated)

Joystick disabled, all dental chair movement controls are locked.

Protection against liquid penetration.

The foot control is protected against liquid penetration.
Degree of protection: IPX1.

Cleaning.

Clean the foot control using a suitable product (see Paragraph 1.4).



5.2.5. WIRELESS FOOT CONTROL

The "side travel" and "push-pedal" foot controls can also be supplied in WIRELESS version. The wireless foot control contains a ZIGBEE transmitter module (module certified for Europe, Canada and the USA).

Warnings for use.

- WARNING:**
 - Avoid keeping the WIRELESS foot control in proximity of other RF sources, such as wireless LAN cards, other radio devices, home RF devices, microwave ovens. The recommended distance is at least 2 metres in the case of microwave ovens and 1 metre in all other cases.
 - Even though the electromagnetic field irradiated by the foot control is insignificant, it is advisable NOT to use it in proximity of life support equipment (e.g. pacemakers or heart stimulators) and hearing aids. Before using any electronic device in health facilities, always check that it is compatible with the other equipment present.
 - Exclusively use the dental unit to charge the battery of the WIRELESS foot control.
 - The internal battery may only be replaced by a qualified technician.

Warnings for first use.

It is advisable to fully charge the foot control battery before using it for the first time.

WIRELESS foot control operation.

The WIRELESS foot control operates exactly in the same way as the wired version, therefore refer to the paragraphs above paying WARNING to the specific model used.

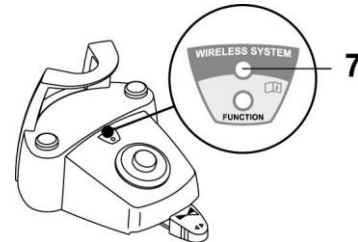
In addition, the WIRELESS foot control has a specific LED (7) that indicates the battery charge and the communication status with the dental unit.

LED (7) indications.

The colour of the LED indicates the battery charge, while the type of flashing indicates the communication status with the dental unit.

Battery charge:

COLOUR	DESCRIPTION (CABLE DISCONNECTED)	DESCRIPTION (CABLE CONNECTED)
GREEN	Battery charge (>75%)	Battery charged
ORANGE	Battery charge (<50%)	Battery charging
RED	Battery needs charging (<25%)	Battery charge error
Off	Battery flat	Dental unit off or foot control fault



Communication status:

FLASHING	DESCRIPTION
Slow	Connection active in wireless mode
Fast	Connection active with charging cable inserted
Double	Connection search
On fixed	Communication error

Battery characteristics.

The WIRELESS foot control is equipped with a rechargeable Lithium-Polymer battery (Li-Poly, 3.7V, 5200 mAh type Guangzhou Markyn Battery Co. Model 9051109).

The battery life is approximately 2 months (estimating 8 hours of consecutive daily operation) with the battery fully charged and fully efficient. The battery efficiency reduces with age. It is estimated that the efficiency is reduced to 60% after 500 complete recharging cycles. Also in this condition, the battery should last about 1 month.

NOTE: when the battery efficiency is so reduced as to be deemed unsatisfactory to support the daily usage requirements, have it replaced by a qualified technician (original spare part no. 97901336).

WARNING: Do not attempt to replace the battery yourself.



Limited battery warranty.

The battery in the foot control is covered by a 6-month warranty from the date of installation.

Recharging the battery.

When the batteries in the WIRELESS foot control need to be recharged, Proceed as follows:

- Open the protective cap of the connector on the rear of the foot control and connect the recharging cable.
- Connect the other end of the recharging cable to the dental unit (see figure).

At this point, the foot control is in the battery charging phase (battery charging warning LED on) even though remaining fully functional.



NOTE:

the battery is fully recharged in about 6 hours.



WARNING:

Exclusively use the dental unit to charge the battery of the WIRELESS foot control.

Natural battery discharge.

Should the battery not be used for long periods of time, it may slowly discharge all the same.

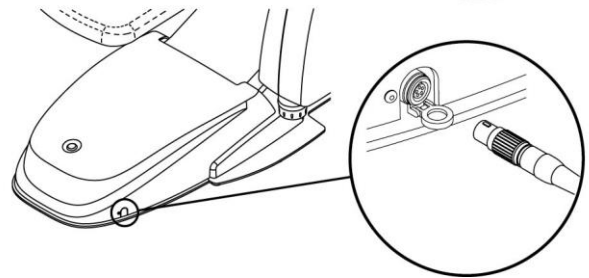
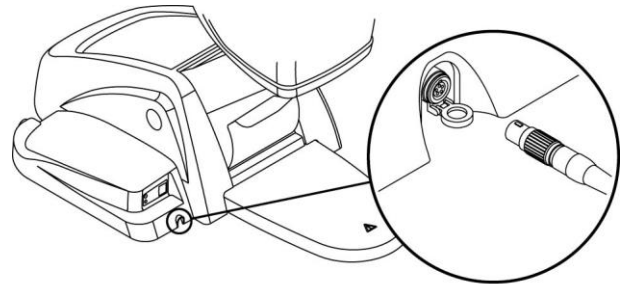
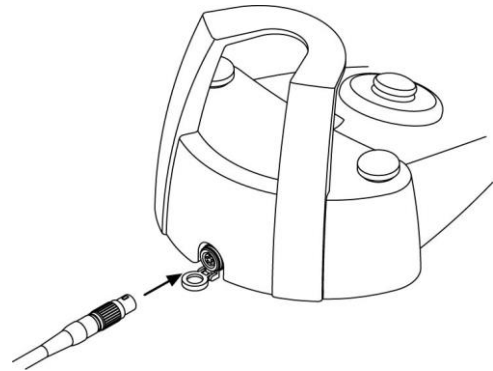
If unused for long periods of time, it is advisable to always fully charge the battery before use.

Maintenance and disposal

The wireless foot control does not contain parts that can be repaired directly by the user.

In the event of a malfunction, do not attempt to carry out maintenance operations, but directly contact the manufacturer or his local distributor at the numbers indicated in the warranty certificate.

At the end of its lifetime, the battery must be replaced by a specialised technician at a Service Centre.





5.3. SYRINGE

Description of the instrument.

- a Nozzle.
- b Handpiece.
- c Syringe release button.
- d Air button.
- e Water button.
- f Hot/cold selector (only for 6-function syringes).
- g Hot/cold indicator LED (only for 6-function syringes).

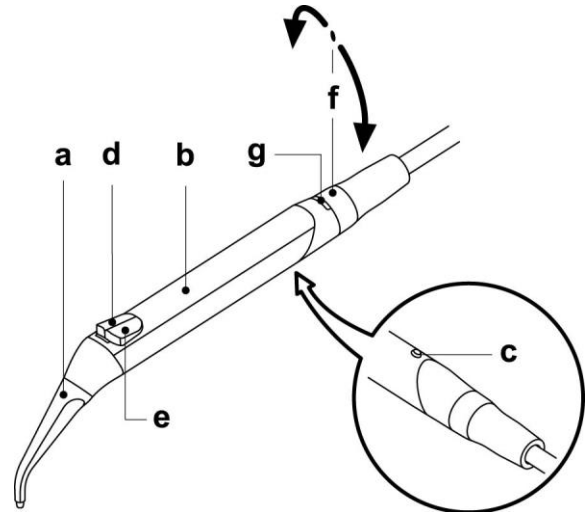


WARNING:

The instrument is supplied non-sterile.
It is recommended to use disposable protections and nozzles.

Technical specifications.

- Operating time:
 - syringe 3F: continuous operation,
 - syringe 6F: 5 sec. operation, 10 sec. rest,
- Power supply:
 - 6F syringe (CEFLA s.c. models): 24 Vac; 50/60 Hz; 2 A; 50 W.
- Classification in accordance with standard EN 60601-1:
 - 6F syringe (CEFLA s.c. models): CLASS II, type B
- consult the Technical Installation Manual (see Paragraph 11.).



Use.

3-function syringes:

- Place the instrument in its working position.
- Button (e) = water;
- Button (d) = air;
- Button (e + d) = spray.

6-function syringes:

- Place the instrument in its working position.
- Operation with hot spray, air and water: turn the selector switch (f) clockwise (LED g on).
- Operation with cold spray, air and water: turn the selector switch (f) counter-clockwise (LED g off).
- Button (e) = water;
- Button (d) = air;
- Button (e + d) = spray.

Removing the grip.

- The nozzle (a) is screwed onto the grip (b).
- 3-function syringes:
 - Press the button (c) to take the grip off the syringe casing.
- 6-function syringes:
 - Turn the selector switch counter-clockwise (LED g off) and press the button (c) to take the grip off the syringe casing.

Syringe removable cord.

Syringe features a removable cord to make cleaning operations easier (see paragraph 5).

Cleaning.

Use soft disposable paper towel dampened with detergents/disinfectants.



WARNING:

- Do not soak the syringe in disinfectant liquids or detergents.
- Recommended products: harsh products and/or products containing acetone, chlorine and sodium hypochlorites.

Disinfection.

Syringe grip and nozzle in a steam autoclave at 135°C (2 bar) following the instructions for use of the device.



NOTE:

bag before sterilising.



5.4. TURBINE

Connecting the handpiece and changing the drill.

Refer to the specific instructions provided with the handpiece.

Use.



WARNING:

Read the instructions for use of the various turbines.

- **Operating times: work 5 min., rest 5 min.**
- Place the instrument in its working position.
- Use the foot control lever to start the instrument (see paragraph 5.2.).
- The cock (f), found near the instrument, is used to adjust the amount of water in the spray.
- The cock (e) adjusts the amount of air spray for all the instruments.



NOTE:

the turbine cord can also be used to connect the air micromotors equipped with 4-way connector and conforming to ISO 13294 Standard - Dental Air Motor.

Adjusting the turbine's operating speed.

- Place the instrument in its working position.
- Use buttons INCREASE or DECREASE to set the maximum speed percentage of the turbine.



NOTE:

data settings are automatically saved.

- The foot control pedal adjusts the operating speed from the minimum to the maximum values set (see paragraph 5.2).

Turning on the fiber optics.

- Place the instrument in its working position.
- Press the button to turn the Fiber Optics on/off.



NOTE:

the Fiber Optics switch off when the turbine is not used for 30 seconds (rheostat switch off).

Syringe removable cord.

Syringe features a removable cord to make cleaning operations easier (see paragraph 5).

Cleaning and care.

Refer to the specific instructions provided with the handpiece.

It is recommended to use Daily Oil (CEFLA s.c.) for lubrication.

Disinfection.

Steam in autoclave at 135°C (2 bar) following the instructions for use of the device.



WARNING:

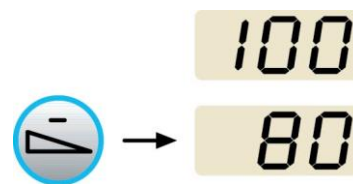
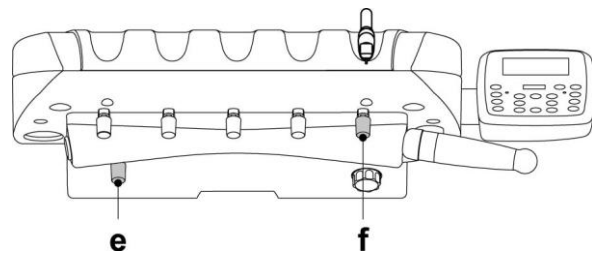
Carefully read the operating instructions supplied with the handpiece before attempting to sterilise.

Warnings for use.



WARNING:

- The instrument is supplied **NOT STERILE** and must be sterilized before use.
Carefully read the operating instructions supplied with the instrument before attempting to sterilise.
- The turbine must never be started without attaching the drill or fake drill.
- The drill release button must not be pressed down during operation!
Friction between button and micromotor impeller overheats the head and may cause burns.
- The patient's internal tissues (tongue, cheeks, lips, etc.) must be protected against contact with the button by using suitable instruments (mirrors, etc.).
- The drills and various instruments attached to the handpieces must comply with Biocompatibility Standard ISO 10993.





5.5. MICROMOTOR

Installing the handpieces and changing the drill.

Refer to the specific instructions provided with the micromotor and various handpieces.

Use.



WARNING:

Also read the instructions for use of the various motors.

- **Operating times: work 5 min., rest 5 min.**
- Place the instrument in its working position.
- Use the foot control lever to start the instrument (see paragraph 5.2.).
- The cock (**f**), found near the instrument, is used to adjust the amount of water in the spray.
- The cock (**e**) adjusts the amount of air spray for all the instruments.

Adjusting the micromotor's operating speed.

- Place the instrument in its working position.
- Use the INCREASE or DECREASE buttons to set the maximum micromotor rotation speed.



NOTE:

data settings are automatically saved.

- The foot control pedal adjusts the operating speed from the minimum to the maximum values set (see paragraph 5.2).

Selecting the direction of rotation.

- Place the instrument in its working position.
- Press the INVERSION button on the console.
LED off: clockwise rotation.
LED on: counter-clockwise rotation.



WARNING:

When the micromotor is taken out, 3 BEEPS sound to signal counter-clockwise rotation.



NOTE:

when the rheostat lever is on, the micromotor cannot reverse its direction of rotation.

Turning on the fiber optics.

- Place the instrument in its working position.
- Press the button to turn the Fiber Optics on/off.



NOTE:

the fiber optics switch off when the micromotor is not used for 30 seconds (rheostat switch off).

Syringe removable cord.

Syringe features a removable cord to make cleaning operations easier (see paragraph 5).

Cleaning and care.

Refer to the specific instructions provided with the handpiece.

It is recommended to use Daily Oil (CEFLA s.c.) for lubrication.



WARNING:

- **Do not soak the syringe in disinfectant liquids or detergents.**
- **Recommended products: harsh products and/or products containing acetone, chlorine and sodium hypochlorites.**

Disinfection.

Handsets only: steam autoclave at 135°C (2 bar) following the instructions for the use of the device.



WARNING:

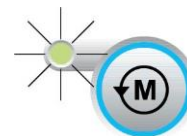
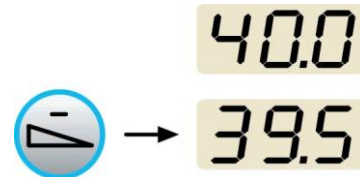
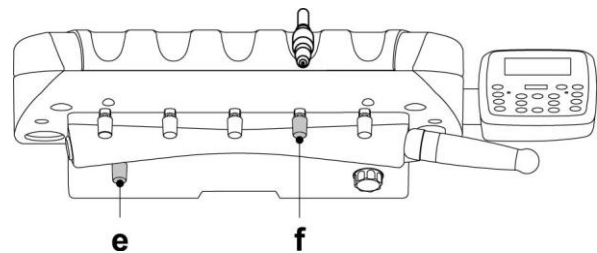
Carefully read the operating instructions supplied with the instrument before attempting to sterilise.

Warnings for use.



WARNING:

- **The instrument is supplied NOT STERILE and must be sterilized before use.**
Carefully read the operating instructions supplied with the instrument before attempting to sterilise.






- Never put the contra angle on the micromotor while it is running.
- The drill release button must not be pressed down during operation!
Friction between button and micromotor impeller overheats the head and may cause burns.
- The patient's internal tissues (tongue, cheeks, lips, etc.) must be protected against contact with the button by using suitable instruments (mirrors, etc.).
- The drills and various instruments attached to the handpieces must comply with Biocompatibility Standard ISO 10993.

5.6. SCALER

Connecting the handpiece and insert.

Refer to the specific instructions provided with the handpiece.

-  **WARNING:**
Before attempting to connect the handpiece, make sure the contacts are perfectly dry. Blow air from the syringe, if necessary, to dry.

Use.


-  **WARNING:**

Also read the instructions for use of the various handpieces.

- Operating times: see operating instructions supplied with the handpiece.
- Place the instrument in its working position.
- Use the foot control lever to start the instrument (see paragraph 5.2).
- The cock (f), found near the instrument, is used to adjust the cooling water flow.

Adjusting the scaler's power output.

- Place the instrument in its working position.
- Use the "INCREASE" or "DECREASE" buttons to set the scaler's maximum power percentage.


-  **NOTE:**
the data set are automatically saved.

- The foot control pedal operates the instrument at the set maximum power (see paragraph 5.2).

ENDO function.

The scaler operates at up to ½ of the maximum power set.

- Place the instrument in its working position.
- Press the INVERSION button on the console.
LED off: standard operation.
LED on: ENDO function active.

-  **NOTE:**
with the foot control lever activated, you cannot change operation.

Syringe removable cord.

Syringe features a removable cord to make cleaning operations easier (see paragraph 5).


Cleaning and care.

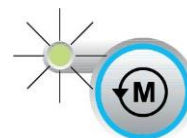
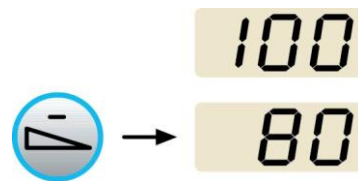
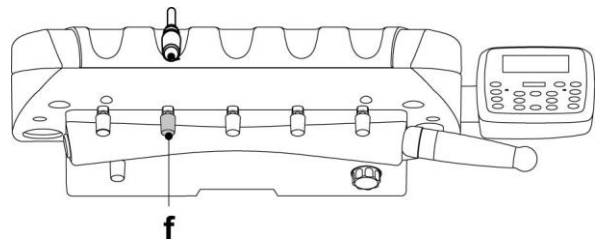
Refer to the specific instructions provided with the instrument.

-  **WARNING:**
- Do not soak the handpiece in disinfectant liquids or detergents.

Disinfection.

- Torque wrench, scaler bits and scaler handset: steam autoclave at 135°C (2 bar) following the instructions for the use of the device.

-  **WARNING:**
Carefully read the operating instructions supplied with the instrument before attempting to sterilise.



**Warnings for use.****WARNING:**

- The instrument is supplied **NOT STERILE** and must be sterilised before use.
Carefully read the operating instructions supplied with the instrument before attempting to sterilise.
- Before attempting to connect the handpiece, make sure the contacts are perfectly dry.
Blow air from the syringe, if necessary, to dry.
- Make sure the threaded sections of the insert and handpiece are perfectly clean.
- Do not change the shape of the insert.
- Check wear and tear of the insert on a regular basis, replacing it in the following cases:
 - obvious wear,
 - drop in performance,
 - deformation or impact.
- **Notes for PIEZOLIGHT 6 scalers:**
 - Class 1 LED apparatus;
 - Do not direct the light beam in anyone's eyes when cleaning or servicing the device (it is recommended to keep the fiber optics switched off).
- To avoid hazards or malfunctions when connecting the board, do not reverse the positions of the cords for scalers of different brands.
- The inserts fitted on the handpiece must comply with ISO 10993 Standard on biocompatibility.



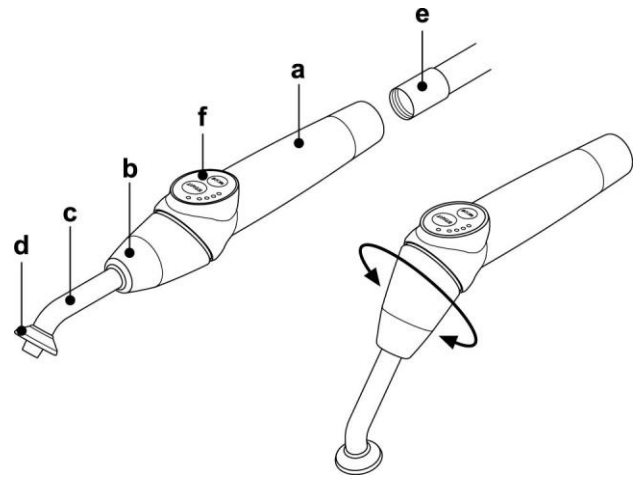
5.7. T LED CURING LIGHT

Technical specifications.

Supply voltage: 24-36 VDC
 Max. power absorbed: 6 VA
 Light source: 1 5W LED
 Wavelength: 430-490 nm
 Acoustic signals: at cycle start, every 5 seconds, and at cycle end
 Type of operation: intermittent (work 3 consecutive cycles - rest 60 sec.)
 Programs: 6 (preset).

General description of the lamp.

- a Lamp handpiece.
- b Rotary end.
- c Fiber optic.
- d Eye protection.
- e Power cord.
- f Control console.



NOTE:

the curing light can be used in different configurations (wand-style, gun-style or any intermediate position) as it is more convenient for the user.



NOTE:

the curing light is delivered in its original packing which should be kept for future shipment.

Description of the control pad.

1 LED 1 (STANDARD cycle) :

Emission of 1000 mW/cu.cm for 20 seconds (this cycle is set as default at the time of sale).

2 LED 2 (FAST cycle) :

Emission of 1600 mW/cu.cm for 15 seconds.

3 LED 3 (STRONG cycle) :

Emission of 1800 mW/cu.cm for 20 seconds.

4 LED S :

When LED S is on, you access ramp cycle mode and at the same time the LEDs B, R and L next to it come on:

[LED S + LED 1] ramp cycle B (BONDING) :

Ramp cycle with emission of 500 mW/cu.cm for 5 seconds, ramp from 500 to 1000 mW/cu.cm for 5 seconds and 1000 mW/cu.cm for 5 seconds for a total of 15 seconds.

[LED S + LED 2] ramp cycle R (RAPID RESTORATION) :

Ramp cycle with emission of 500 mW/cu.cm for 5 seconds, ramp from 500 to 2200 mW/cu.cm for 5 seconds and 2200 mW/cu.cm for 5 seconds for a total of 15 seconds.

[LED S + LED 3] ramp cycle L (LONG RESTORATION) :

Ramp cycle with emission of 500 mW/cu.cm for 5 seconds, ramp from 500 to 1800 mW/cu.cm for 5 seconds and 1800 mW/cu.cm for 10 seconds for a total of 20 seconds.

5 Malfunction signalling LED :

This red LED comes on only if there is a malfunction.

6 START button :

Pressing the START button starts the cycle currently selected (the cycle LED will come on).

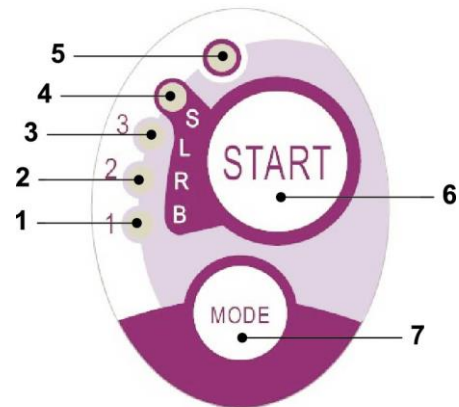
If it is pressed again at any time during the cycle, light beam emission will immediately be interrupted.

7 MODE button :

This button is used to select the cycle to be run. It allows changing from the current cycle to the immediately following cycle.

The first three cycles (1, 2 and 3) are at constant power and the LEDs come on individually. When LED S is on, you access ramp cycle mode and at the same time the LEDs B, R and L come on.

Once the LED of the cycle you intend to use has come on, the lamp is ready for use. Pressing the START button, light beam emission is activated according to the selected cycle.



Cycle	LED	Total time	Ø8 mm	Total energy
STANDARD	1	20"	1,000 mW/cm ²	20,000 mJ
FAST	2	15"	1,600 mW/cm ²	24,000 mJ
STRONG	3	20"	1,800 mW/cm ²	36,000 mJ
BONDING	S+1	15"	ramp cycle	11,250 mJ
RAPID REST.	S+2	15"	ramp cycle	20,250 mJ
LONG REST.	S+3	20"	ramp cycle	26,250 mJ




NOTE:

the cycle can be selected and the button is operative only when the curing light is not emitting any light. If the button is accidentally pressed while light is being emitted, nothing will happen.

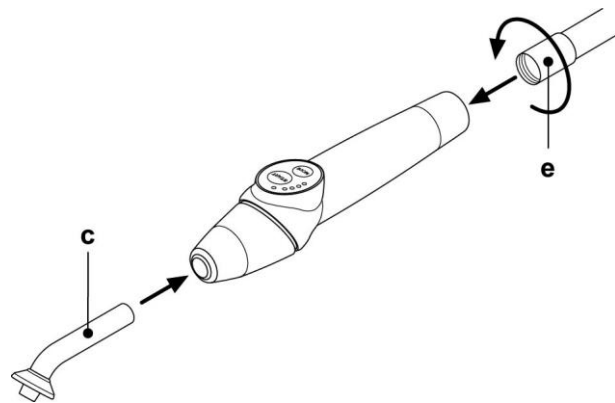



Operation.

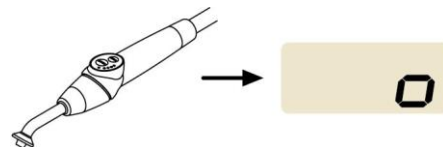
WARNING:
 **The instrument is supplied non-sterile.**

Before use, disinfect the lamp grip. The optical fibre and the eye protection can be sterilised in a steam autoclave at 135°C.


- Put the fiber optics (**c**) in its housing until it clicks.
- Attach the curing light handpiece to the end of its power cord and tighten the ring nut (**e**).
- Take the lamp out of its housing on the assistant's board or dentist's board.




NOTE:
 *one animated symbol will be shown on the console display to confirm that the instrument has been activated.*




- Turn the front of the lamp and/or fiber optics to the position most suitable for curing (wand, gun or intermediate position).
- Use the MODE button to select the desired cycle as previously directed (the selected cycle is always indicated by the illuminated LED).


NOTE:
 *the curing light has a permanent memory therefore the last cycle used will always be present the next time it is used.*

- Place the fiber optics in the position required for curing.


NOTE:
 *the fiber optics should be placed as close as possible to the material to be cured without touching it.*

- Press button START to start the cycle.

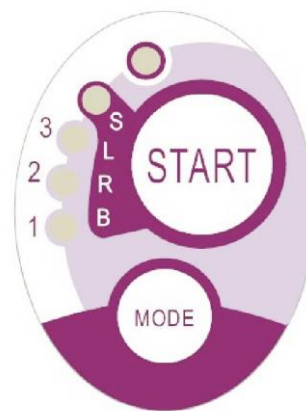
WARNING:
 **Operating mode: work 2 consecutive cycles, rest 60 seconds.**

NOTE:
 *when a programmed cycle is activated, the LEDs (1, 2, 3, B, R, L) indicate the time that elapses (in multiples of 5 seconds) and turn off every 5 seconds of operation. The curing light comes with a beep that BEEPS when the cycle starts, BEEPS every 5 seconds of operation and lastly BEEPS twice at the end of the work cycle.*

- Allow light emission to stop by itself. However, it can be stopped at any time by simply pressing the START button again.

WARNING:


- The curing light is equipped with a system that signals malfunctions by illuminating the LEDs in different combinations (see next paragraph).
- The curing light is equipped with a cut-out.




Indicators.

The following indicators are provided on the control console to signal curing light fault:

- **LED 5 and LED 1, green, steady on.**
Lamp does not emit any light.
Contact technical service department.
- **LED 5 and LED 2, green, steady on.**
Instrument start up microcontroller fault.
Contact technical service department.
- **LED 5 and LED 3, green, steady on.**
Power supply too low.
Contact technical service department.
- **LED 5 and LED 4 flash continuously.**
Handpiece cut-out tripped. These LEDs will continue to flash until the light has cooled down enough (about 5 minutes) to be used again.
If the problem persists, contact technical service department.

Maximum curing thickness.

The maximum curing thickness with every single cycle is 3 millimetres (refer to the instructions of the composite material used as well).

WARNING:
 **This thickness must not be exceeded as the layer may not be completely cured.**

**Warnings for use.****WARNING:**

The LED is a Class 2 light source in accordance with IEC 62471 standard. DO NOT STARE AT THE BEAM.

The light emitted may cause eye injury in case of direct radiation without eye protection.

Always use an eye protection shield when operating the curing light and do not direct the light beam to the eyes.

The light emitted may damage soft tissues (oral cavity mucous membrane, gums, skin).

Be extremely careful to aim the light precisely on the material to be cured.

- **People with eye diseases, such as those who have had cataracts removed or retina diseases must be adequately protected when the curing light is used, for example with suitable protective eyewear.**
- **The rotary end can turn 180° counter-clockwise in relation to the handpiece to change from wand to gun configuration.**
To go back to wand configuration, turn clockwise.
A click is heard when these two positions are reached. Do not turn any more once the click is heard.
The intermediate positions can be used even if a click is not heard.
Put the fiber optics back into the correct position after turning the end section.
- **Do not pull the power cord.**
- **Do not expose the handpiece to excessive vibrations.**
- **Do not drop the handpiece and in particular the fiber optics.**
The lamp may break if accidentally banged.
Check the condition of the handpiece if it has been banged or dropped before using the curing light.
Try to turn on the light and check operation first without using it on the patient.
If cracked, broken or if there are any other faults, do not use the curing light on the patient and contact the technical service department.
The fiber optics is rather delicate and may crack or break if banged, affecting the final amount of light emitted.
If dropped, carefully inspect the fiber optics to verify if it is cracked or broken. If cracked, a strong light appears in the spot in which the fiber is cracked.
In all these cases, the fiber optics must be replaced.
- **The curing light handpiece (sold separately) can be connected only to dental units with connections for this curing light handpiece. Connection to any other equipment may damage the circuits inside the lamp and seriously injure the user and patient.**
- **The curing lamp handpiece is not protected against liquid penetration.**
- **The curing lamp handpiece is not suitable for use in the presence of flammable anaesthetic gas mixed with air, oxygen or nitrous oxide (N₂O).**

Cleaning.

The curing light may be a means of cross contamination between patients.

The most contaminated parts are the fiber optics and the eye protection. Before sterilising them, make sure there are no residues of curing products: if necessary, clean with alcohol or a plastic spatula.

Exclusively sterilise the optical fibre and the eye protection in an autoclave at a sterilisation temperature of at least 134°C.

**WARNING:**

- **The fiber optics can undergo 500 autoclave cycles after which it tends to become opaque and therefore emit less light.**
- **The eye protection must also be replaced after 500 cycles.**
- **Contact the manufacturer to purchase original spare parts (fiber optics + eye protection: code 97660404).**

The handpiece cannot be put in autoclave; disinfect it on the outside with suitable products and cover it with disposable plastic wrap.

Use soft disposable paper towels to disinfect the handpiece. Do not use harsh products or soak in liquids.

**WARNING:**

- **The curing light handpiece is NOT suitable for autoclave.**
- **The curing light handpiece is not protected against penetration of liquids therefore it CANNOT be soaked in a solution to be sterilized.**
- **The outside of the lamp should be disinfected with the fiber optics on.**
Do not use any type of disinfectant on the exposed optical surface of the handpiece when the fiber is removed. The surface will become irreparably opaque if it comes into contact with disinfectant.

Maintenance.

This equipment does not require any particular type of maintenance.

Only technicians authorised by the manufacturer can replace and/or repair the handpiece and dental unit.

The handpiece has been purposely constructed in a manner that requires specific tools to open it and therefore it cannot be removed by the user.

The warranty is automatically void if the handpiece is altered in any way.

Troubleshooting.

- **When the lamp is removed, the light does not come on (no LED on control console illuminated).**
Make sure the Midwest connection is correctly attached to the power cord.
Carefully screw the ring nut, try to put the lamp back in and then take it out again.
If the problem persists, contact technical service department.
- **Less light emitted.**
 - Make sure the fiber optics is not cracked or damaged in any way: replace it if it is so.
Contact the manufacturer to purchase original spare parts.
 - Make sure there are no residues of curing products on the end of the fiber optics: if necessary, wipe off with alcohol or a plastic spatula.

If the handpiece must be sent back, please disinfect it.

Ship it back in its original packing.

In addition, please enclose a description of the fault with the shipping note.



5.8. C-U2 DENTAL CAMERA

The C-U2 dental camera system, complete with an extremely lightweight ergonomic handpiece, is specially designed for simple and well-conceived usability in examining the oral cavity. Auto-exposure and fixed focus features provide easy operation. This system is designed to allow the dentist to more efficiently show and explain to patients all oral conditions and reasons for planned treatment. The C-U2 system allows filming and taking high-definition (1280x720) live images of the section in question to be taken through a touch of a fingertip on the touch-sensitive area of the handpiece. The live intraoral images are displayed on the monitor or Personal Computer.



WARNING:

The camera may be used as a tool to aid in diagnosis; however, the result must always be supported with visual examination and/or other diagnostic indications.

Evaluations and conclusions based only on the image taken by the camera may be poor as the colours and shapes, electronically processed, may not perfectly correspond to the actual ones.

Warnings for use.



WARNING:

- The external PC and the external monitor must be of medical grade, namely they have to be certified and comply with the standard IEC 60601-1 3rd Ed. They have to be able to ensure a double insulation level for both patient (2 MOPP) and operator (2 MOOP):
 - with respect to the power mains;
 - to all the I/O ports (USB, LAN) supplied with Safety Extra Low Voltage (SELV).
- Even though the electromagnetic field irradiated by the device is insignificant, it is advisable not to use it in proximity of life support equipment (e.g. pacemakers or heart stimulators) according to the specifications included in the user manual of such equipment.
- The disposable infection control sheaths must be used with the device. Change the sheath for each new patient.
- After putting on a new disposable infection control sheath, check it over before using the camera, making sure it is not torn anywhere. If it is, take it off and put on a new one.
- Do not place the handpiece in liquids or in autoclave under any circumstances.
- Store the handpiece in a clean dry area.
- Do not bend the connecting cable excessively.
- Be extremely careful not to drop the handpiece and do not expose it to excessive vibrations.
- Never use a damaged handpiece. Make sure the camera is in good condition and has no sharp edges before attempting to use it. If in doubt, do not use the handpiece, carefully put it away, and contact technical assistance.
- Before starting the equipment, check the condition of the lens protection.
- Do not aim the light beam at the operator's or patient's eyes during operation.
- During continuous use (example, more than 10 consecutive minutes), the temperature of the camera's tip usually increases significantly; if this is uncomfortable, put the handpiece in its holder for a few minutes to allow the light source to cool down. When the camera needs to be used for a prolonged time, reduce light brightness.
- If left running for extended periods, make sure the temperature of the tip is acceptable before attempting to use the camera. Briefly touch the clear plastic part with your fingertip being careful not to touch the lens in the middle.
- Do not attempt to bend, pull or remove the handpiece.

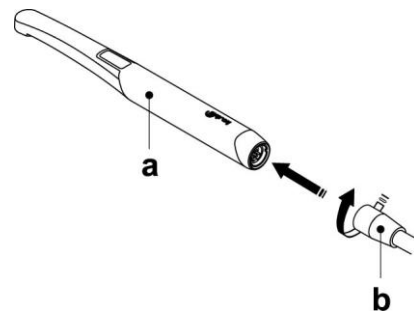
Connecting the handpiece.

Attach the handpiece of camera C-U2 (a) to the end of the cord and tighten the ring nut (b).



WARNING:

Make sure the cord is firmly screwed onto the handpiece.



Use of the camera.

Place the handpiece in its work position; the light comes on and the camera is in the last LIVE mode used.



NOTE:

two animated symbols will be shown on the console display to confirm that the instrument has been activated.

Turning on camera lighting system.

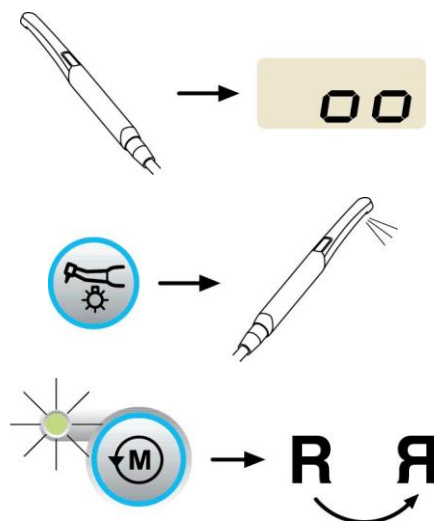
Press the FIBER OPTICS button to turn on/off camera lighting system.

MIRROR function.

With the camera set to LIVE mode, briefly press the INVERSION button present on the console to change from viewing real images to mirror images.

LED off: real viewing.

LED on: mirror viewing.





• FREEZE function.

Handpiece C-U2 is equipped with a touch-sensitive button (g). Tap this button or actuate the foot control to freeze the image of interest. Tap this button again (or actuate the foot control) to unlock the frozen image.

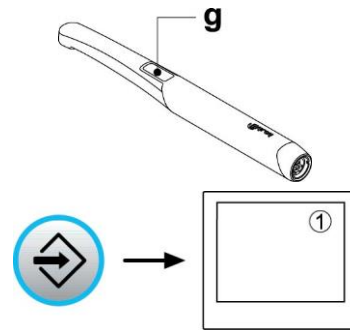
• Setting operation in single image or multiple image mode

With the camera set to LIVE mode, briefly press the SAVE button present on the console to change from viewing a single image to multi-images (and vice versa).



NOTE:

an icon (1) found in the top right-hand corner signals when this mode has been selected.



• Operation in single image mode.

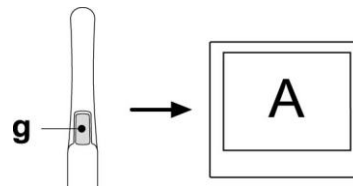
The "live" image appears on the monitor when the camera is set to LIVE and single image mode.

Pressing the touch-activated button (g) on the handpiece (or activating the foot control), you can freeze the image, which is immediately shown on the display deleting any other previous image.



NOTE:

the last image frozen remains on the monitor even if the camera is put back in place.



• Operation in multiple image mode.

- The "live" image appears on the monitor and an icon (1) is shown in the top right-hand corner when the camera is set to LIVE and multi-image mode.

Pressing the touch-activated button (g) on the handpiece (or activating the foot control), you can freeze the image, which is immediately shown on the display.



NOTE:

the frozen image appears directly on the monitor in the first box available on the page currently displayed. Any following frozen image is then positioned in the following box in reading order. Once the 4 available boxes have been filled, each following frozen image will replace the existing ones appearing always in reading order.

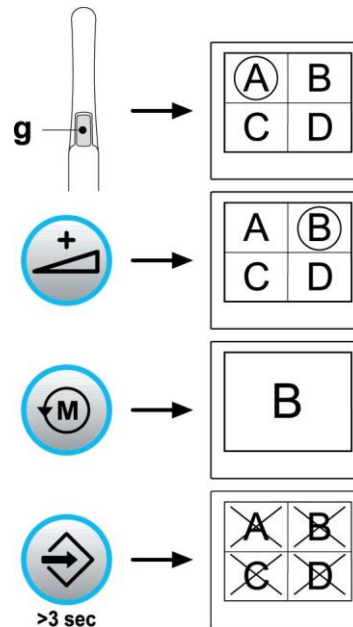
- With the camera in FREEZE mode, pressing the INCREASE or DECREASE buttons or activating the CHIP-AIR command using the foot control (see paragraph 5.2), you can select the 4 stored images in rotation.



NOTE:

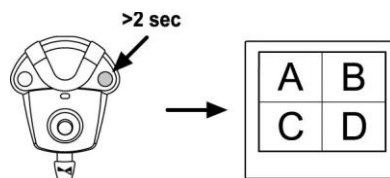
press the SAVE button for 3 seconds to delete all 4 images on the page displayed.

- With the camera in FREEZE mode, pressing the INVERSION button or activating the WATER CLEAN SYSTEM command using the foot control (see paragraph 5.2), you can activate/deactivate full-screen viewing of the selected image.



• VIEW function.

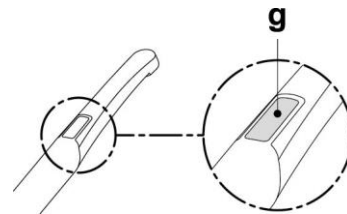
With the camera in LIVE mode, pressing the WATER CLEAN SYSTEM button on the foot control for at least 2 seconds (see paragraph 5.2), you can view the previously frozen images without storing new ones.





Handpiece status.

An optical guide, illuminated by a multicolour LED indicator, found in the area near the control button (g), shows handpiece status as per the table given below:



Colour	Situation
Blue light flashes, very slowly	Handpiece in standby
Light blue steady light	Handpiece activated, live images displayed
Blue/ light blue flashing light	Handpiece in image freeze mode
Brief red flashes	Internal error: contact Technical Service

MyRay iCapture.

This programme allows the C-U2 camera to be set up when it is connected to a PC/WORKSTATION.

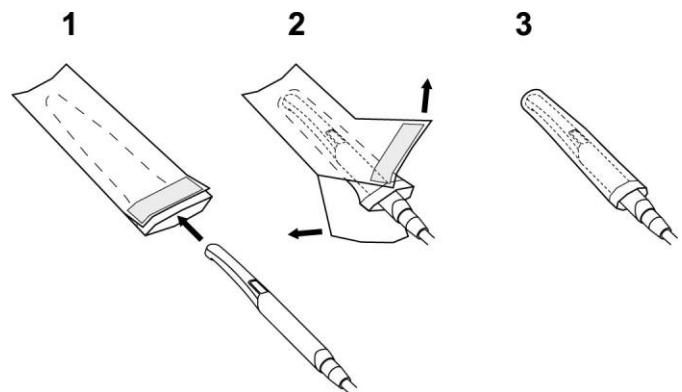
For a full description on how the MyRay iCapture programme works, refer to the instructions, in electronic format, supplied with the C-U2 handpiece.



Disposable infection control sheaths.

The camera can be a source of cross-contamination between patients. For this reason **always use it with a disposable infection control sheath** (code 97901590) and disinfect it on the outside after use everyday. The sheath (with white paper backing) is enclosed in two protective layers: a transparent one with blue tab at the front and a paper one at the back. Follow the directions below to install a new infection control sheath:

- 1 Insert the camera handpiece tip between the layer with White tab and the rear paper backing. The lens, surrounded by the LEDS, must face down, towards the rear paper layer. Gently push the handpiece to the end of the sheath.
- 2 Pull the blue tab removing the protective films.
- 3 The camera is now protected and ready for use.



WARNING:

- Always make certain the handpiece is correctly inserted inside the infection control sheath.
- Always change the disposable infection control sheath before using the device on a new patient.
- Disposal: the disposable infection control sheaths are to be treated as special waste materials (like surgical gloves).

Cleaning and disinfection.

Clean the handpiece with a suitable product after each use: refer to paragraph 1.4.



WARNING:

- The dental camera is not designed for cold sterilization by being soaked, for example, in solutions such as glutaraldehyde or hydrogen peroxide.
- All products must be used as directed by the manufacturer.
- All materials used to clean and disinfect must be thrown away.

Maintenance and repairs.

The C-U2 dental camera does not require any particular maintenance. In the event of malfunctions, please send back the complete handpiece.



WARNING:

There are no parts that can be repaired on site. In the event of a malfunction, please contact an authorized dealer.

Returning parts.

- Please send back any defective devices in their original packaging. Do not reuse damaged boxes.
- The device must be disinfected before being shipped to prevent cross-contamination. Handpieces that have not been adequately cleaned and disinfected will not be accepted.



WARNING:

The sender shall be held responsible for any equipment damage occurred during shipment regardless of whether or not the devices are under warranty.

6. ASSISTANT'S BOARD OPERATION

Dental units model PUMA ELI R - PUMA ELI R CP - PUMA ELI R CART - PUMA ELI R ORTHO:

(1) Assistant's board with twin arm.

Main features:

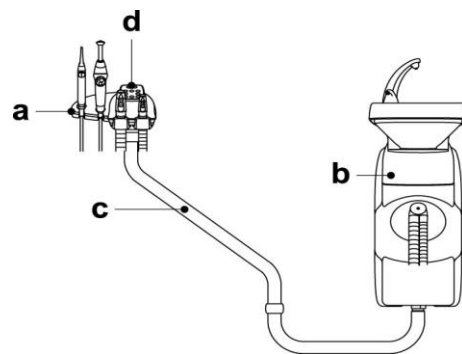
- Two articulated arms (c) secure the board (a) to the water unit (b) allowing it to be placed in the most convenient work position.
- The assistant's board (a) comes with a control console (d) with buttons used to operate the dental chair and water unit.
- The assistant's board can hold 2 suction tubes and 2 instruments.
- The assistant's board comes with sliding rollers (f) that guide and hold up the suction tubes.



NOTE:

the assistant's board is equipped with a safety device that locks out dental chair movement when the board itself is pushed upwards.

(1)



(2) Assistant's board with single arm.

Main features:

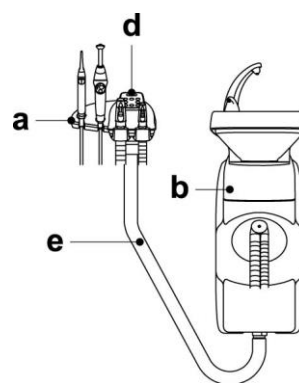
- A single articulated arm (e) secures the board (a) to the water unit (b) allowing for movement within a wide range of action.
- The assistant's board (a) comes with a control console (d) with buttons used to operate the dental chair and water unit.
- The assistant's board can hold 2 suction tubes and 2 instruments.
- The assistant's board comes with sliding rollers (f) that guide and hold up the suction tubes.



NOTE:

the assistant's board is equipped with a safety device that locks out dental chair movement when the board itself is pushed upwards.

(2)



Dental units model PUMA ELI A R and PUMA ELI A R CP:

(3) Assistant's board with 3 arms.

Main features:

- Three articulated arms (c) secure the board (a) to the water unit (b) allowing it to be placed in the most convenient work position.
- The pantograph arm (g) allows a vertical stroke of the assistant's board of 335 mm, in 5 working positions.



NOTE:

to take assistant's board back to bottom position, press button (h) on pantograph arm.

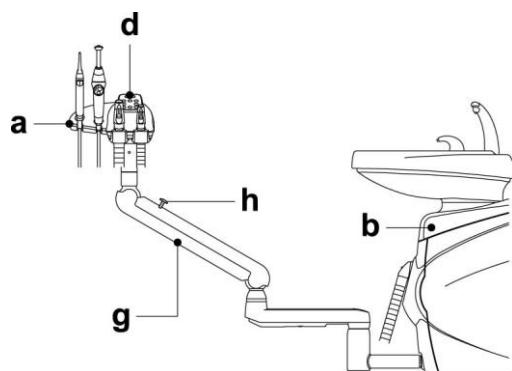
- The assistant's board (a) comes with a control console (d) with buttons used to operate the dental chair and water unit.
- The assistant's board can hold 2 suction tubes and 2 instruments.
- The assistant's board comes with sliding rollers (f) that guide and hold up the suction tubes.



NOTE:

the assistant's board is equipped with a safety device that locks out dental chair movement when the board itself is pushed upwards.

(3)



(4) Assistant's board with single arm.

Main features:

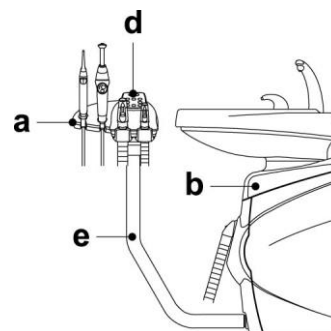
- A single articulated arm (e) secures the board (a) to the water unit (b) allowing for movement within a wide range of action.
- The assistant's board (a) comes with a control console (d) with buttons used to operate the dental chair and water unit.
- The assistant's board can hold 2 suction tubes and 2 instruments.
- The assistant's board comes with sliding rollers (f) that guide and hold up the suction tubes.



NOTE:

the assistant's board is equipped with a safety device that locks out dental chair movement when the board itself is pushed upwards.

(4)

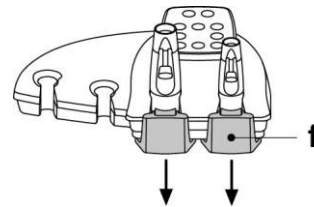




Cleaning the sliding rollers.

Push down and take off the sliding rollers (f).

Clean the sliding rollers with a suitable product: refer to paragraph 1.4.



6.1. ASSISTANT'S BOARD CONSOLE

Description of the buttons:



Water-to-bowl button.



Water-to-cup button.



Operating light on/off button.



Hygiene procedure activation button.



Automatic return position recall button.



Rinsing position recall button.



Chair seat up button.



Chair backrest up button.



Chair seat down button.

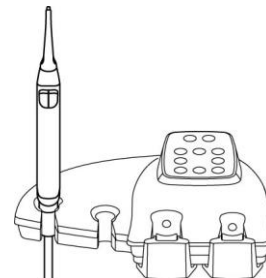


Chair backrest down button.



6.2. SYRINGE

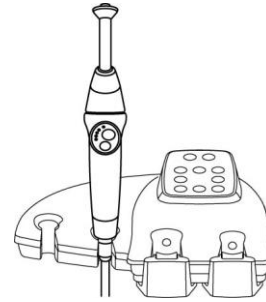
For detailed information regarding operation of this instrument, see paragraph 5.3.





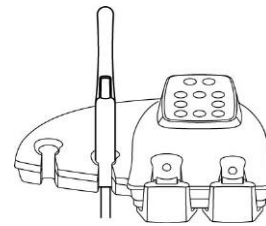
6.3. T LED CURING LIGHT

For detailed information regarding operation of this instrument, see paragraph 5.7.



6.4. C-U2 DENTAL CAMERA

For detailed information regarding operation of this instrument, see paragraph 5.8.

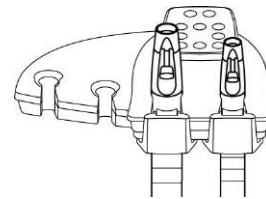


6.5. SUCTION TUBES

Suction starts by taking the tube off the board.
To adjust suction power, use the slider (a) located on the tube holder grip.



NOTE:
when the tube is put back in place, suction stops approximately 2 seconds later. This is done to dry the suction tubes.



Removing the suction tubes.



WARNING:
Always wear goggles and gloves to prevent contact with infected material when removing the suction tubes.

Remove the suction tubes from the conveyor fittings by turning and pulling the tube fitting.
Detach the suction tubes from the holders by turning and pulling the tube fitting.



WARNING:
Never directly grasp the suction tube.

Flushing the suction tubes.

As the dental units may be equipped with different suction systems (liquid ring or wet, air) carefully follow the instructions provided by the suction system manufacturer when disinfecting the system regarding the product to be used, times and directions for use.



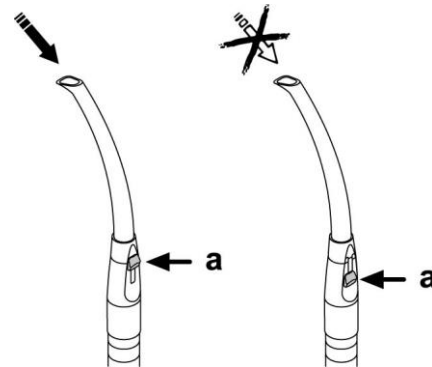
WARNING:
For cleaning of the suction system, it is recommended to use STER 3 PLUS (CEFLA s.c.) diluted in a 6% solution (equivalent to 60 ml of product in 1 litre of water).

Disinfection.

- Cannula-holder terminals: steam autoclave at 135°C (2 bar) following the instructions for the use of the device.
- Suction tubes: soak to cold-sterilise.



WARNING:
With the tubes, never use procedures where temperature goes over 55°C.





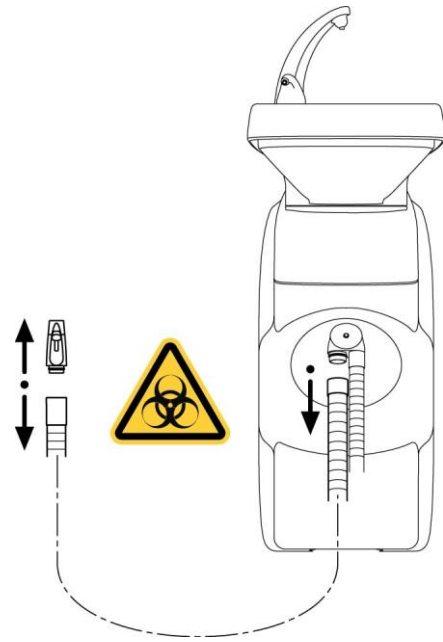
Maintenance.

Periodically lubricate the O-rings of the cannula holder terminals (see Paragraph 9.4.) using **S1-Protective Lubricant for O-Rings** CEFLA s.c..

Note about biocompatibility.

Only use suction tubes supplied with the dental unit and original replacement tubes.

The suction tubes must comply with Biocompatibility standard ISO 10993.



6.6. SALIVA HYDRAULIC SUCTION UNIT

The hydraulic saliva ejector starts running when the tube is removed from the support.

Cleaning after each use.

Aspirate about ½ litre of STER 3 PLUS (CEFLA s.c.) diluted in a 6% solution (equivalent to 60 ml of product in 1 litre of water).

Cleaning the saliva ejector filter.

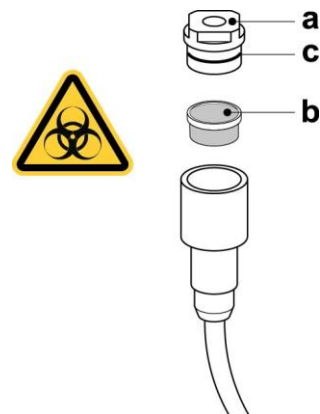
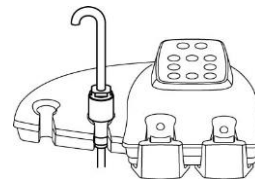
This operation must be carried out at the end of each work day.



WARNING:

Put on goggles and gloves before attempting to perform this operation!

- Suck about ½ litre of STER 3 PLUS diluted in a 6% solution (equivalent to 60 ml of product in 1 litre of water).
- In order to prevent possible dripping of liquids and matter from the filter (**b**) to be extracted, suck only air for about 5 seconds.
- Take off the cap (**a**) by turning and pulling at the same time.
- Remove the filter (**b**).
- Clean/replace the filter (code 97290060).
- Put the filter and cap back in place.



NOTE:

to prevent liquids and matter from dripping from the filter taken out, suck air only for approximately 5 seconds before performing these operations.

Routine maintenance.

Lubricate the O-Ring [**c**] with **S1-Protective Lubricant for O-Rings**.



6.7. TURBINE

For detailed information regarding operation of this instrument, see paragraph 5.4.

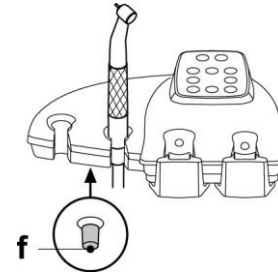
The cock (**f**), found under the board, is used to adjust the amount of water in the spray.



NOTE:

for PUMA ELI R ORTHO model:

as it does not feature any dentist's console, turbine rotation speed cannot be set and any fibre optics is always on.





7. WATER UNIT OPERATION

7.1. FILL CUP AND BOWL

The bowl can be turned 305° on the water unit by hand.
The bowl, the water-to-cup spout and the water-to-bowl spout can be removed to make the cleaning operations easier.

Control buttons.



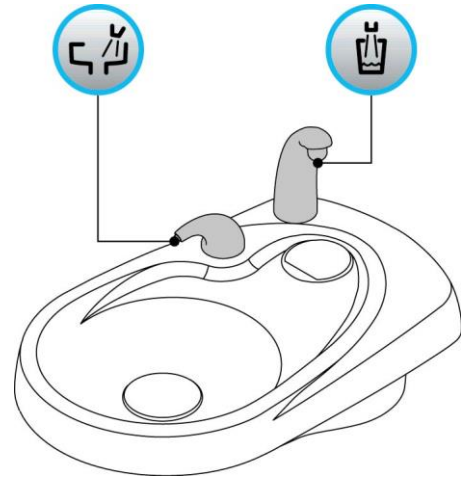
Water-to-cup button.



Water-to-bowl button.

Bowl flushing automatically stops after 30 seconds.
Bowl flushing automatically starts in the following cases:

- when the "Rinse Position" button is pressed;
- when the "Water to cup" button is pressed.



Hot water to cup.

When this function is provided, hot water is always delivered to the cup.

Adjusting the amount of water used to fill the cup.

• Turn on the operating unit holding down the Water to cup button on the assistant's board.



NOTE:

1 intermittent BEEP signals saving is in progress.

• Press the water to cup button once to start filling the cup.
• Once the desired water level has been reached, press the water-to-cup button again making sure the BEEP stops.

At this point, the new water level in the cup has been saved.



Changing bowl operation.

Water delivery to the bowl can take place in an untimed manner (ON/OFF operation with the relative button).

This modification is made by turning on the operating unit holding down the Water-to-Bowl button on the assistant's board.

A beep sounds to signal the selected operating mode.

• 1 BEEP: timed operation.
• 3 BEEPS: ON/OFF operation.



NOTE:

timed operation is factory set.

Changing the water to bowl controller with water delivered to cup.

The controller that delivers water to the bowl when the "water to cup" button is pressed can be enabled/disabled.

This modification is made by turning on the operating unit simultaneously holding down the Water to cup and Water to bowl buttons on the assistant's board.

A beep sounds to signal the selected operating mode.

• 1 BEEP: controller enabled.
• 3 BEEPS: controller disabled.



NOTE:

the controller is enabled by default.



Water-to-bowl controller modification upon recall of dental chair "Rinse position".

The controller that delivers water to the bowl when the dental chair "Rinse Position" button is pressed can be turned on/off.

This modification is made by turning on the operating unit simultaneously holding down the dental chair "Rinsing Position" and "Water-to-Bowl" buttons on the assistant's board.

A beep sounds to signal the selected operating mode.

- 1 BEEP: controller enabled.
- 2 BEEPS: controller disabled.



NOTE:

the controller is enabled by default.

Water-to-bowl controller modification upon recall of dental chair "Automatic return".

The controller that delivers water to the bowl when the dental chair "Automatic Return" button is pressed can be turned on/off.

This modification is made by turning on the operating unit simultaneously holding down the dental chair "Automatic Return" and "Water-to-Bowl" buttons on the assistant's board.

A beep sounds to signal the selected operating mode.

- 1 BEEP: controller enabled.
- 3 BEEPS: controller disabled.



NOTE:

the controller is enabled by default.

Taking off the spouts, bowl and bowl filter.

- Pull up the spouts (l) and (n) and take them off.
- Pull up the filter (q) and its cover (p) from the bowl to remove them.
- Pull up the bowl (m) and take it off.

Disinfecting and cleaning.



WARNING:

Always wear goggles and gloves to prevent contact with infected material when cleaning the bowl and bowl filter.

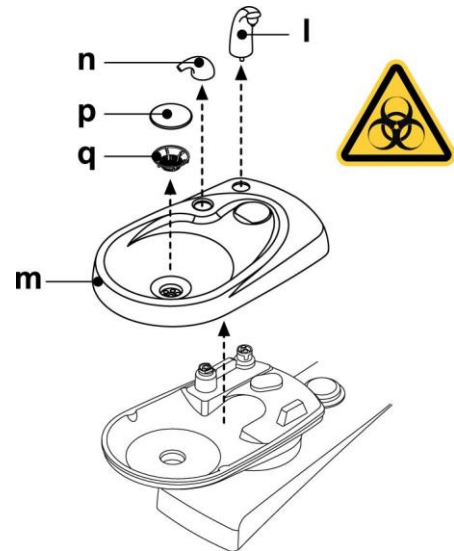
The parts are to be cleaned daily at the end of each working day.

- Water-to-cup and water-to-bowl spouts: sterilise the cannula holder terminals in a steam autoclave at 135°C (2 bar) following the instructions for use of the device.
- Bowl: thoroughly wash with a specially formulated scale-remover.
- Bowl filter: clean with running water and commercially-available cleaning products.



WARNING:

Do not use acid or harsh products.





7.2. S.S.S. SYSTEM. -ST

Applied on models PUMA ELI R , PUMA ELI R CP e PUMA ELI R CART, PUMA ELI A R , PUMA ELI A R CP, only.


Description of the system.

The system S.S.S. -ST (Separate Supply System) features a tank (a) for distilled water.

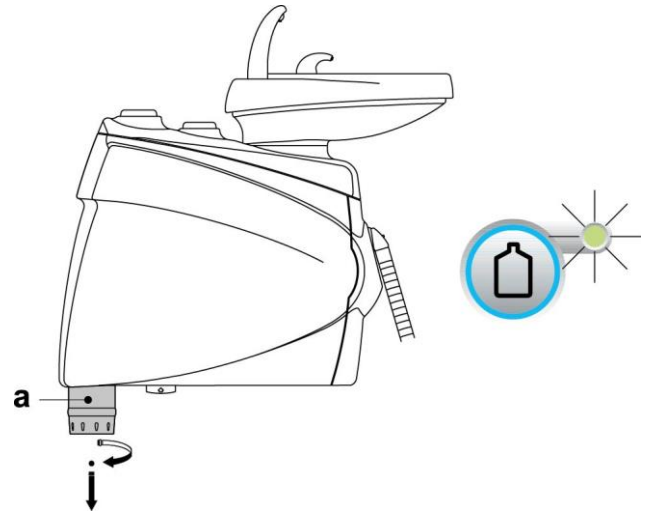
Tank has a total capacity of 1.1 litres.

Tank supplies:

- the sprays of all the instruments present on dentist's and assistant's boards,
- the syringe present on the assistant's board,
- the cup filling system (optional),
- if available, water quick coupling (optional).


Button , present on dentist's board console allows enabling/disabling the S.S.S. -ST system:

- LED off: system not active.
- LED on: system active.



Tank filling.

Once tank min.level is reached (approx. 200 cc), fill it as follows:

- Disable the S.S.S. -ST system by pressing button .



NOTE:

during this operation the pressurised air present inside tank will be automatically bled to the outside.

- Remove tank (a) by turning it clockwise.
- Pour distilled water inside tank until reaching the max. level.



WARNING:


Use distilled water only as, for enhanced hygiene conditions, can be charged with 600 parts per million of hydrogen peroxide using 20 ml of Peroxy Ag+ per litre of distilled water, or oxygenated water (20 ml of 3% oxygenated water per litre of distilled water).

- Fit tank back in place by turning it counter-clockwise.



WARNING:

Make sure that tank is duly fastened.

- Press again button  to enable the S.S.S. -ST system and confirm that it has been duly filled up.



WARNING:

If you are going to be absent from the surgery for long periods of time (holidays), completely empty the tank (a) before leaving.

Tank cleaning.

We recommend cold sterilising tank only at regular intervals (at least once a month) using a peracetic acid based product and proceeding as follows:

- remove tank from dental unit and drain it completely,
- prepare the solution of the peracetic acid based product by following the instructions given by the manufacturer,
- fill tank to the brim with the peracetic acid solution,
- leave the peracetic acid solution inside tank for the time recommended by the manufacturer,
- drain all peracetic acid solution from tank,
- rinse tank with distilled water,
- fill tank with distilled water, if necessary after adding the above-indicated substances,
- fit tank back in place inside dental unit.



7.2.1. S.S.S. SYSTEM. MANUAL

Applied on models PUMA ELI R ORTHO and PUMA ELI R SINGLE CART, only.

Description of the system.

System features a tank (a) for distilled water. Tank has a total capacity of 1.1 litres (models PUMA ELI R ORTHO) and 1.8 litres (models PUMA ELI R SINGLE CART).

Tank supplies:

- the sprays of all the instruments present on dentist's and assistant's boards,
- the cup filling system (optional),
- if available, water quick coupling (optional).

A by-pass toggle (b) allows disabling the system in case you want to supply instruments with mains water.


This system allows performing a disinfection cycle of instrument spray ducts (see paragraph 7.2.2.).

WARNING:
 Carry out a disinfection cycle at the end of every work day.


Tank filling.

Once tank min. level is reached (approx. 200 cc. for models PUMA ELI R ORTHO and approx. 500 cc. for models PUMA ELI R SINGLE CART), fill it as follows:


- Turn the toggle (c) to position "CLOSE AIR PRESSURE".
- Remove tank (a) by turning it clockwise.

NOTE:
 during this operation the pressurised air present inside tank will be automatically bled to the outside.

- Pour distilled water inside tank until reaching the max. level.

WARNING:
 Use distilled water only as, for enhanced hygiene conditions, can be charged with 600 parts per million of hydrogen peroxide using 20 ml of Peroxy Ag+ per litre of distilled water, or oxygenated water (20 ml of 3% oxygenated water per litre of distilled water).

- Fit tank back in place by turning it counter-clockwise.
- Turn the toggle (c) to position "OPEN AIR PRESSURE".

WARNING:
 Make sure that tank is correctly secured in place before turning toggle (c) to position "OPEN AIR PRESSURE".
 • If you are going to be absent from the surgery for long periods of time (holidays), completely empty the tank (a) before leaving.

BY-PASS control.

System can be disabled in case you want to supply instruments with mains water.

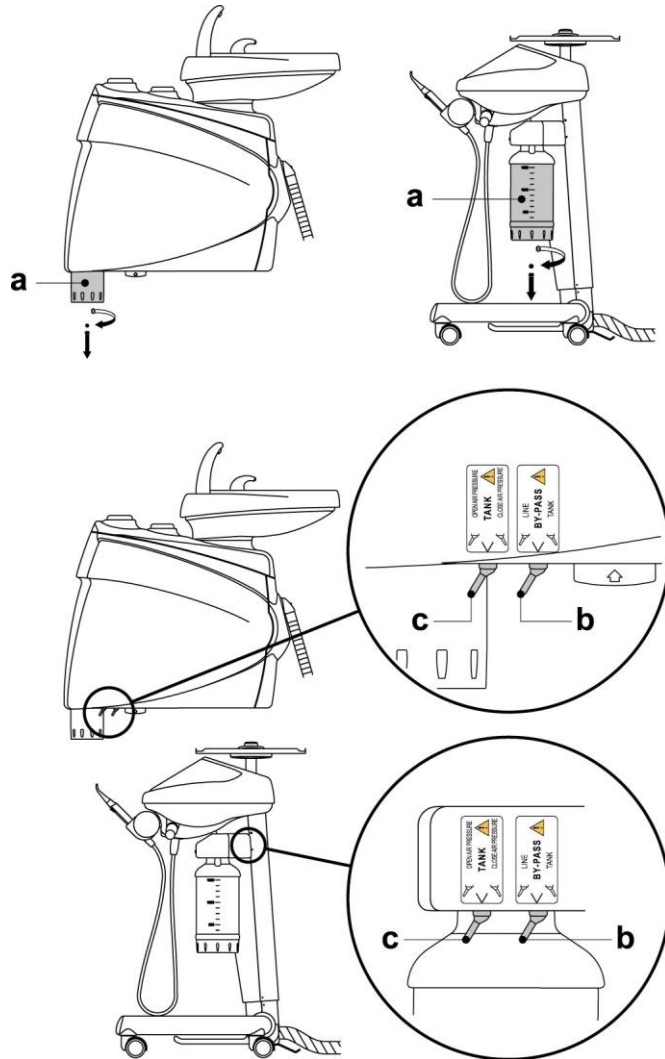
This function is carried out by turning toggle (b) to position "LINE".

Turn toggle (b) back to position "TANK" to go back to distilled water operation.

Tank cleaning.

We recommend cold sterilising tank only at regular intervals (at least once a month) using a peracetic acid based product and proceeding as follows:

- remove tank from dental unit and drain it completely,
- prepare the solution of the peracetic acid based product by following the instructions given by the manufacturer,
- fill tank to the brim with the peracetic acid solution,
- leave the peracetic acid solution inside tank for the time recommended by the manufacturer,
- drain all peracetic acid solution from tank,
- rinse tank with distilled water,
- fill tank with distilled water, if necessary after adding the above-indicated substances,
- fit tank back in place inside dental unit.





7.2.2. MANUAL DISINFECTION CYCLE

The S.S.S. system allows performing a manual disinfection cycle of the water ducts of all the instruments on the dentist's board, on the assistant's board and on the water-to-cup duct (optional).

To disinfect, proceed as directed below:

A) Preparing the disinfectant solution:

- Pour undiluted PEROXY Ag+ (or 3% oxygenated water) into the tank marked with an orange band.



NOTE:

make sure that the tank is completely filled.

B) Disinfectant injection phase:

- Make sure the spray cock of each dynamic instrument is fully open (if it is not, either very little or no water at all will flow out).
- Turn the toggle (c) to position "CLOSE AIR PRESSURE".
- Remove the distilled water bottle (a) by turning it clockwise.
- Replace the bottle of distilled water (a) with the one containing PEROXY Ag+.
- Remove the syringe and empty the duct with the water button.



NOTE:

when emptying is complete, no water will come out of the duct.

- Repeat the same procedure for the syringe on the assistant's board (with the water key). Use the Water clean system with the foot control for the remaining dynamic instruments (see paragraph 5.2).
- If foreseen, drain the water-to-cup duct by pressing the "Water-to-cup" button.
- Turn the toggle (c) to position "OPEN AIR PRESSURE".
- Remove the instruments and fill the ducts with the hydrogen peroxide solution. Use the Water Clean System with the foot control for the dynamic instruments and the water button for the syringes and for the water to cup duct by pressing the "Water-to-cup delivery button".



NOTE:

let the oxygenated water solution flow from the instruments for about 6-7 seconds.

- Put the instruments back in place.



NOTE:

at this point, the ducts contain hydrogen peroxide.

C) Disinfectant contact time:



WARNING:

PEROXY Ag+ must be left to dwell in the ducts for at least 10 minutes, but not more than 30 minutes.

D) Duct rinsing phase:

- Turn the toggle (c) to position "CLOSE AIR PRESSURE".
- Take out the bottle of hydrogen peroxide (a) turning it clockwise.
- Put in the bottle containing distilled water.
- Repeat duct draining procedures for the different instruments and, if necessary, for water-to-cup duct.



NOTE:

when emptying is complete, no water will come out of the duct.

- Turn the toggle (c) to position "OPEN AIR PRESSURE".
- Extract the instruments and let the distilled water flow out using the Water Clean System command of the foot control (see paragraph 5.2) for the dynamic instruments, the water button for the syringes and the "Cup water delivery" button for the cup water delivery duct.



NOTE:

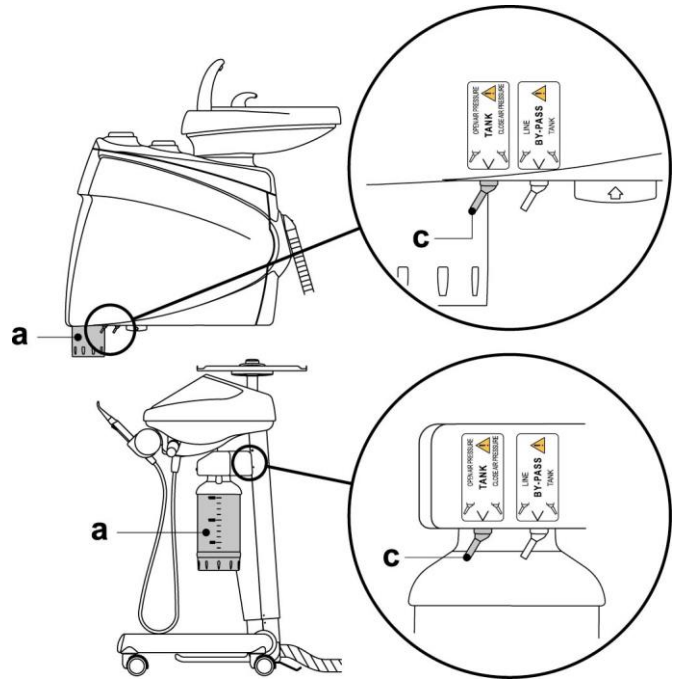
allow approximately 6-7 seconds for water to flow out.

- At this point, the ducts contain distilled water again and the dental unit is ready to be used.



WARNING:

- After disinfection, make sure that you close the tank containing PEROXY Ag+ (exposed to air, it loses its effectiveness).
- It is advisable to run at least one disinfection cycle a day, preferably at the end of the working day.





PEROXY Ag+ storage.

For proper PEROXY Ag+ storage, follow the manufacturer's instructions given on the package.

It is important to keep the package tightly closed and stored in a cool place at a temperature not exceeding 25°C.



WARNING:

Never leave PEROXY Ag+ or oxygenated water in the tank marked with an orange band for more than one month. If you are going to be absent from the surgery for long periods of time (holidays), completely empty the tank marked with an orange band before leaving.

7.3. DISINFECTION SYSTEM

Applied on models PUMA ELI R , PUMA ELI R CP e PUMA ELI R CART, PUMA ELI A R , PUMA ELI A R CP, only.

The AUTOSTERIL -ST system allows performing a disinfection cycle of the water ducts of all the instruments on the dentist's board, of a dynamic instrument located on the assistant's board and of the water-to-cup ducts (optional).

To disinfect, proceed as directed below:

A) Preparing the disinfectant solution:

- Pour undiluted PEROXY Ag+ (or 3% oxygenated water) into the tank marked with an orange band.



NOTE:

make sure that the tank is completely filled.

B) Setting the disinfection cycle:

- Replace tank (a) with the tank filled with disinfectant and activate the S.S.S. -ST system (see paragraph 7.2.).



NOTE:

the AUTOSTERIL -ST cycle can be activated only if the S.S.S. -ST system is enabled.

- Make sure the spray cocks (d) found towards the bottom of the board, are open (if they are not, either very little or no water at all will flow out).
- Position the container (e) for the instruments to be disinfected on the bowl.
- If foreseen, insert the special supplied container (c) under the cup spout.
- Press the "AS" button on the assistant's board for at least two seconds.



NOTE:

an acoustic signal (4 BEEPS) will confirm that control has been acknowledged.

Now the dental unit will enter the stand-by mode, highlighted on the display by several flashing dashes.



NOTE:

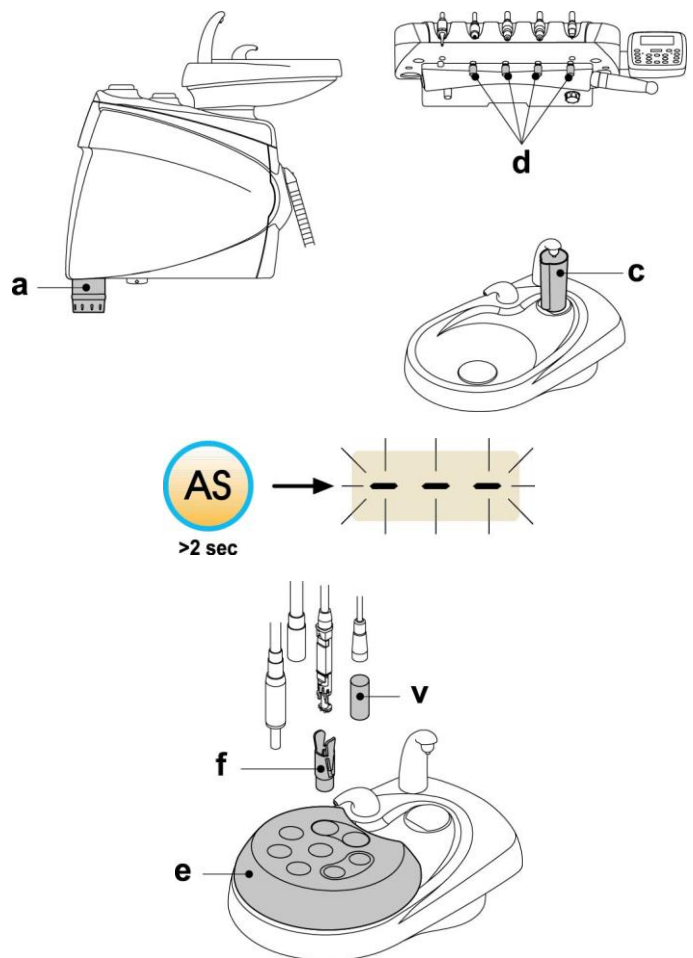
to quit this condition without doing anything, simply keep the "AS" button pressed again for at least 2 seconds.

- Insert the cords of the instruments to be disinfected in the container.



WARNING:

For the syringe instrument you need to use the special adapter (f) and the heating system must be off. Descaler cord must be inserted by using the special adapter (v). The micromotor cord must be fully inserted inside motor body. Turbine cord must be inserted without the handpiece. The syringe positioned on the assistant's board cannot be disinfected through the AUTOSTERIL -ST cycle.





C) Performing the disinfection cycle:

- Shortly press the "AS" button to activate the disinfection cycle.



NOTE:

an acoustic signal (1 BEEP) will confirm that cycle has started.

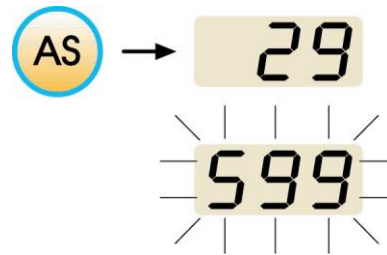
- The system now replaces the water in the ducts.



NOTE:

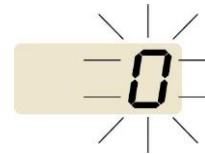
the time remaining to the end of the operation is shown on the console display (duration 30 seconds).

- At the end of water duct filling phase, an acoustic signal (1 BEEP) will be emitted and the time remaining to the end of the disinfectant contact phase will be shown on the console display with flashing figures.



D) Disinfectant contact time:

- Pre-set and not editable contact time: 600 seconds.
- At the end of the disinfectant contact phase, an intermittent acoustic signal (1 BEEP every second) will be emitted for 1 minute. The console display will show that time is now zero and continues flashing.



E) Duct rinsing phase:

- Replace tank (a) filled with disinfectant with the original tank containing distilled water, and activate the S.S.S. -ST system (see paragraph 7.2.).
- Start duct rinsing phase by pressing the "AS" button on the assistant's board.



NOTE:

an acoustic signal (1 BEEP) will confirm that rinsing cycle has started.

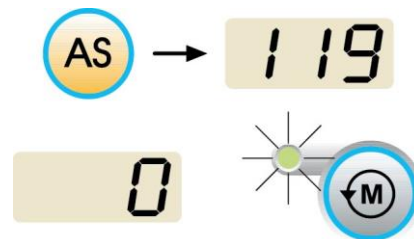
- Now system will replace the disinfectant present inside ducts with distilled water.



NOTE:

the time remaining to the end of the operation is shown on the console display (duration 120 seconds).

- At the end of the rinsing cycle an acoustic signal (1 BEEP) will be emitted, the console will show that time has elapsed (figure 0) and the "INVERSION" button LED will flash.
- At this point, put the instruments extracted back into place to return to the working condition.



WARNING:

- After disinfection, make sure that you close the tank containing PEROXY Ag+ (exposed to air, it loses its effectiveness).
- It is advisable to run at least one disinfection cycle a day, preferably at the end of the working day.

Disinfection cycle interruption.

During setup, you can quit the disinfection cycle by pressing at any time the "AS" button for at least 2 seconds.



NOTE:

once activated, cycle CANNOT be interrupted.

PEROXY Ag+ storage.

For proper PEROXY Ag+ storage, follow the manufacturer's instructions given on the package.

It is important to keep the package tightly closed and stored in a cool place at a temperature not exceeding 25°C.



WARNING:

- Never leave PEROXY Ag+ or oxygenated water in the tank marked with an orange band for more than one month.
- If you are going to be absent from the surgery for long periods of time (holidays), completely empty the tank marked with an orange band before leaving.

7.4. AUTOMATIC INSTRUMENT FLUSHING CYCLE

Description of the system.

The automatic TIME FLUSHING cycle allows to carry out an automatic flushing cycle to renew water present in the water ducts of the instruments on the dentist's and the assistant's boards and the water-to-cup duct.

The flushing cycle can be carried out with mains water or with distilled water (if the S.S.S. -ST or the S.S.S. systems are present)

The cycle time is about 2 minutes.

WARNING:
It is advisable to carry out a flushing cycle at the beginning of each working day and between two patients.

Flushing cycle setting.

• If you want to run a flushing cycle with distilled water, enable the S.S.S. system (see paragraph 7.2. or 7.2.1. based on the available PUMA ELI R model).

NOTE:
it is advisable to execute the flushing cycle with tank (a) completely full.

• Press the "AS" button present on assistant's board to access the TIME FLUSHING mode.

NOTE:
three flashing dashes will be shown on the console display to confirm that TIME FLUSHING mode has been activated.

- Make sure the spray cocks (d) found towards the bottom of the board, are open (if they are not, either very little or no water at all will flow out).
- Position the container (e) for the instruments to be disinfected on the bowl.
- Insert the special supplied container (c) under the cup spout.
- Insert the cords of the instruments to be disinfected in the container.

WARNING:
For the syringe instrument you need to use the special adapter (f) and the heating system must be off. Descaler cord must be inserted by using the special adapter (v).
The micromotor cord must be fully inserted inside motor body. Turbine cord must be inserted without the handpiece.

NOTE:
the syringe is always active, therefore as soon as it is inserted in the instrument container it will immediately start delivering water.

Performing the flushing cycle.

• Start the flushing cycle by pressing the "AS" button on the assistant's console a second time.

NOTE:
the time remaining to the end of the flushing cycle is shown on the console display (duration 120 seconds).

- At the end of the flushing cycle (the value 0 is shown on the display), the LEDs of the FIBRE OPTICS and INVERSION buttons start flashing.
- At this point, put the instruments extracted back into place to return to the working condition.

Flushing cycle interruption.

Once the flushing cycle has been started, it CANNOT be interrupted.

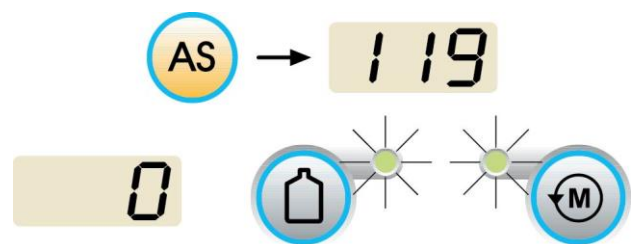
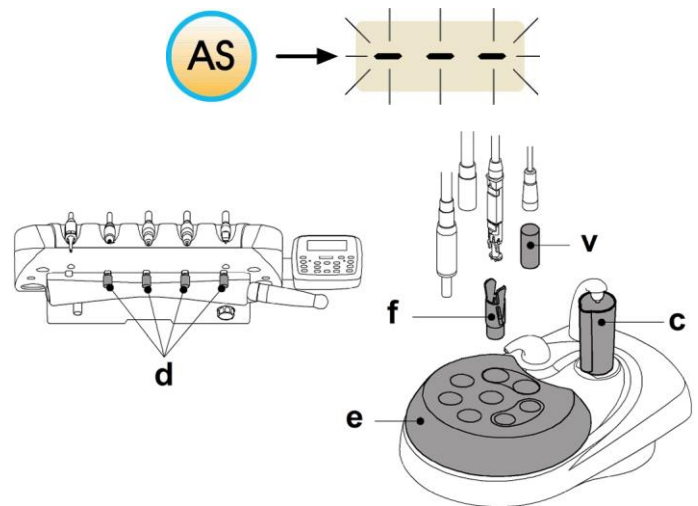
Error signalling.

Signalling: 2 BEEPS when you attempt to start the flushing cycle.

Cause: no instrument extracted or instruments not equipped with water ducts extracted.

Remedy: check the instruments extracted and then start the flushing cycle

NOTE:
if the error message appears again, call Technical Service.

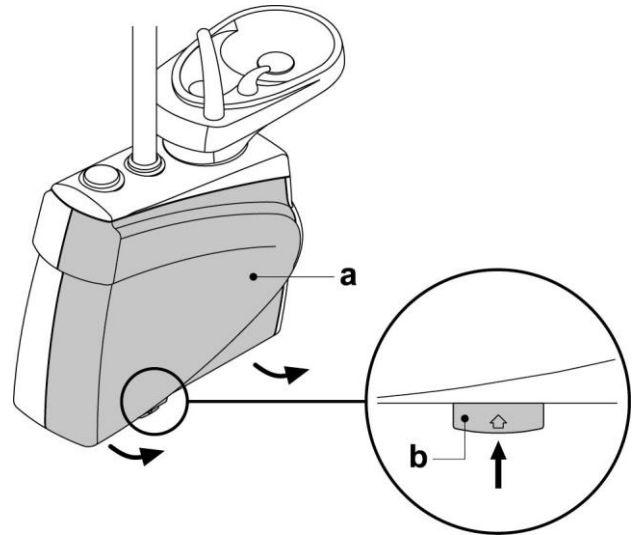




7.5. OPENING/CLOSING THE WATER UNIT SIDE COVER

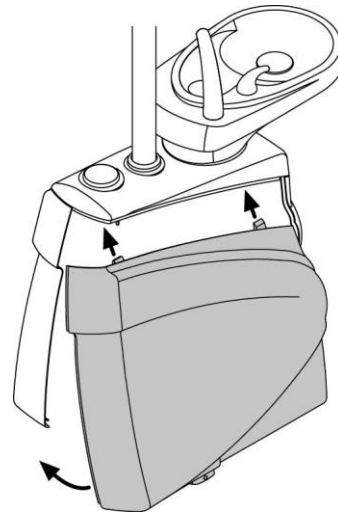
Opening the cover.

- open the cover on the side of the water unit (**a**) after pushing up and releasing the lock lever (**b**).



Closing the cover.

- put on the cover making sure the two locks are inserted in the notches in the water unit,
- lastly, bring the bottom of the cover near the water unit frame to engage the lock lever again.



8. ACCESSORIES

8.1. OPERATING LAMP

The operating light comes in 4 models:

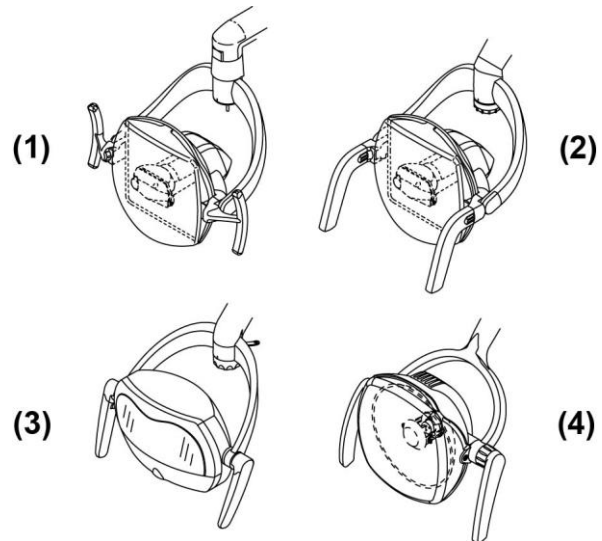
- (1) Light with halogen light source, VENUS model.
- (2) Lamp with halogen light source, model VENUS PLUS.
- (3) Lamp with LED light source, VENUS PLUS-L model.
- (4) Lamp with halogen light source, model VENUS e.

The instructions for use and maintenance of the lamps are available in PDF format and can be downloaded from the download area of the website www.castellini.com.



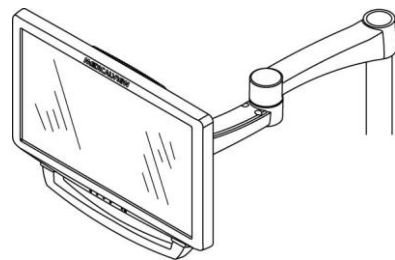
NOTE:

during the automatic movements of the dental chair, the light automatically turns off to prevent blinding the patient.



8.2. MONITOR ON LAMP POLE

The instructions for use and maintenance of the monitor are included with the equipment.



8.3. AUXILIARY TRAY HOLDER

Applied on models PUMA ELI R CART and PUMA ELI R ORTHO, only.
Tray holder can contain two standard size trays.

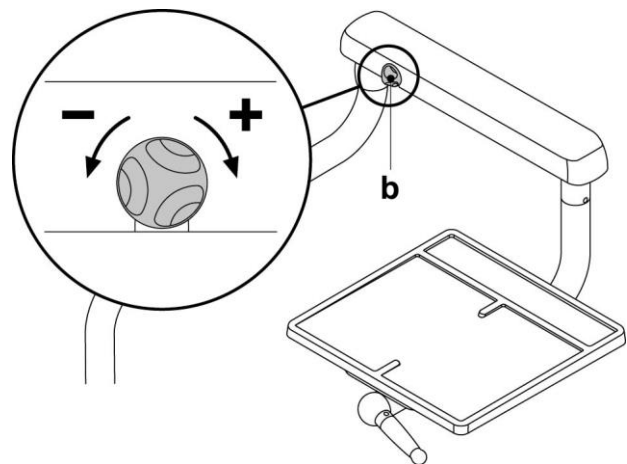
Turn knob (**b**) to adjust vertical movement based on load:

- turn clockwise to increase resistance (heavy loads).
- turn counter-clockwise to decrease resistance (lightweight loads).



WARNING:



Maximum permitted load on the tray: 3.5 Kg (without negatoscope) or 2.5 Kg (with negatoscope).

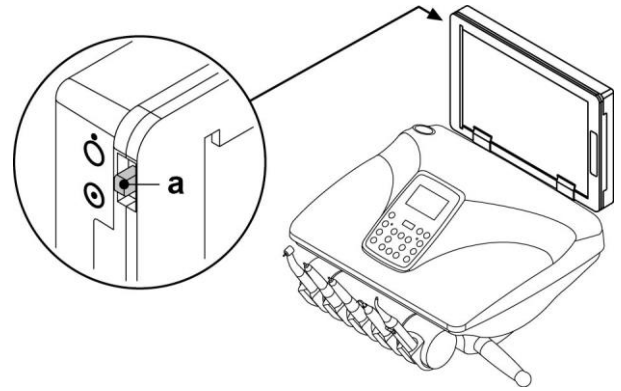




8.4. NEGATOSCOPE FOR PANORAMIC X-RAYS


An x-ray film viewer for panoramic x-rays can be mounted on all INTERNATIONAL version dentist's boards.
The screen dimensions are as follows: H=210mm, L=300mm.
Turn on negatoscope, simply turn special switch (**a**):

-  Negatoscope on.
-  Negatoscope off.




8.5. AIR/WATER/230V QUICK-COUPLINGS

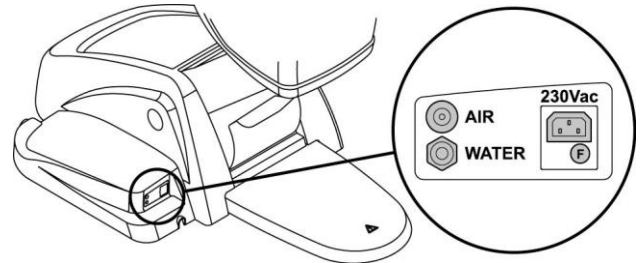
The 230V air/water quick-couplings are located on the side of the utility service centre.

 **WARNING:**
Switch off the equipment before attempting to connect or disconnect the air/water outlets.

Technical specifications.

- Power outlet: 230Vac 2A in accordance with IEC/EN 60320-2-2/F (only on dental units with 230Vac power supply).
- Air quick-coupling pressure: 6 Bar.
- Water quick-coupling pressure:
 - mains water, 2.5 Bar
 - with S.S.S. system, 3 Bar
- Water quick-coupling flow rate:
 - mains water, 1800 ml/min
 - with S.S.S. system, 950 ml/min

 **NOTE:**
with the S.S.S. system, in order to use the quick-coupling with mains water, the distilled water tank needs to be disabled (see paragraph 7.2. or 7.2.1).





9. MAINTENANCE

Preventive maintenance

CEFLA s.c., the manufacturer of the dental units, in accordance with applicable standards IEC 60601-1 3rd Ed. - 2007, IEC 62353 and directive MDD 93/42, and subsequent changes, for medical devices underlines that the preventive maintenance checks for the dental unit specified in the Technical care manual and Maintenance and warranty handbook are to be carried out by authorised personnel at least once every 12 months.



WARNING:

The warranty is void if the equipment is serviced, repaired, altered or modified in any way by personnel who have not been duly authorised by CEFLA s.c..

Safety checks.

In accordance with standard IEC 62353, the safety checks specified in the Technical care manual and Maintenance and warranty handbook supplied with the dental unit are to be carried out at the intervals required by current local regulations. If no precise indications are given, CEFLA s.c., the manufacturer of the dental units, recommends checking them at least every 24 months at the time of installation and whenever electrical parts that are live are repaired/updated.



WARNING:

The manufacturer shall not be held liable for any personal injury or equipment damage if the precautions given above are not observed.

9.1. MAINTAINING THE INSTRUMENTS

Maintenance instructions for the instruments are enclosed with each instrument.



WARNING:

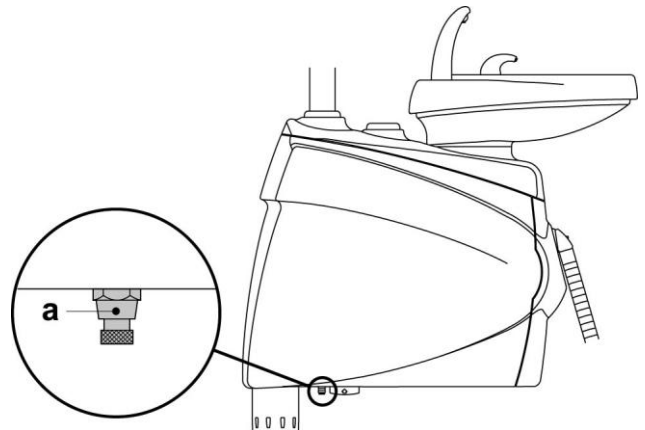
Maintenance of the instruments should be carried out with the equipment switched off.

9.2. DRAINING CONDENSATE

Perform this operation every day before starting work.

Proceed as follows:

- put a container under the cock (a) found below the water unit,
- loosen the cock knob,
- after the tank has been emptied, fully close the cock.





9.3. CLEANING THE SURGICAL SUCTION FILTER

This operation should be done daily at the end of work.

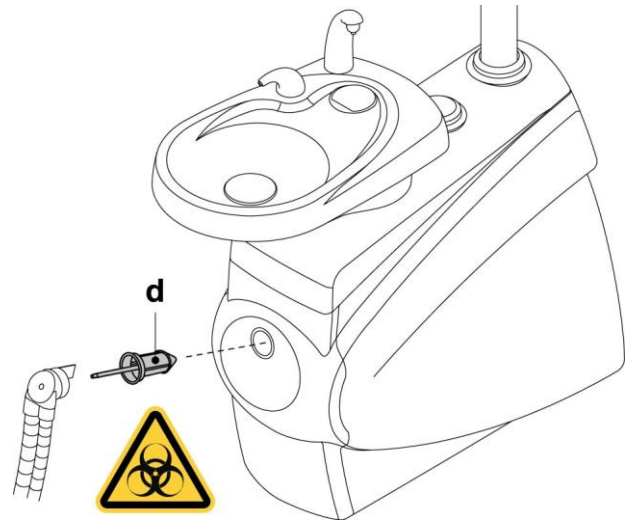
WARNING: Always wear goggles and gloves to prevent contact with infected material when cleaning the suction filters.

Proceed as follows:

- One at a time, remove filter (d).
- Clean/replace the filter (code 97461845).
- Refit filter.

WARNING: Before refitting the filter, make sure to remove any amalgam residues still present on filter seat mouth.

NOTE: in order to prevent any dripping of liquids and secretions from the filter being removed, it is advisable to carry out the above operations with the cannula working.



9.4. SURGICAL SUCTION

The surgical suction system must be sanitized using a product suitable for this purpose.

WARNING: For cleaning of the suction system, it is recommended to use STER 3 PLUS (CEFLA s.c.) diluted in a 6% solution (equivalent to 60 ml of product in 1 litre of water).

At the end of each surgical procedure.

- Suck in about half a litre of solution prepared with the selected disinfectant with each of the suction tubes used.
- Sterilise the cannula holder terminals in a steam autoclave at 135°C (2 bar) following the instructions for use of the device.

At the end of each work day.

- Suck in 1 litre of water with each suction tube, alternating water and air (keep the suction tube alternately in and out of the water).
- Once rinsed with water, suck in approximately half a litre of the solution prepared with the selected disinfectant with each of the suction tubes used.

WARNING: Pay strict attention to all the disinfecting product manufacturer's instructions, warnings and cautions.

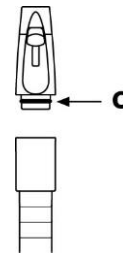
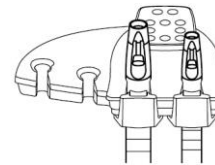
NOTE: after disinfecting, it is good practice to suck in air only to dry the entire suction system (5 minutes).

Once a week.

Remove the cannula body from its cord attachment and lubricate the O-rings (o) using S1-Protective Lubricant for O-Rings (CEFLA s.c.).

Once a year.

Replace the suction tubes and ends of the tube holder.





9.5. CATTANI SURGICAL SEPARATOR

At the beginning of each work day.

Insert inside filter (d) a tablet (v) of VF CONTROL PLUS (CEFLA s.c.)



WARNING:

Always wear gloves to prevent contact with infected material when carrying out this operation.

At the end of each surgical procedure.

- Suck in about half a litre of solution prepared with the selected disinfectant with each of the suction tubes used.
- Sterilise the cannula holder terminals in a steam autoclave at 135°C (2 bar) following the instructions for use of the device.

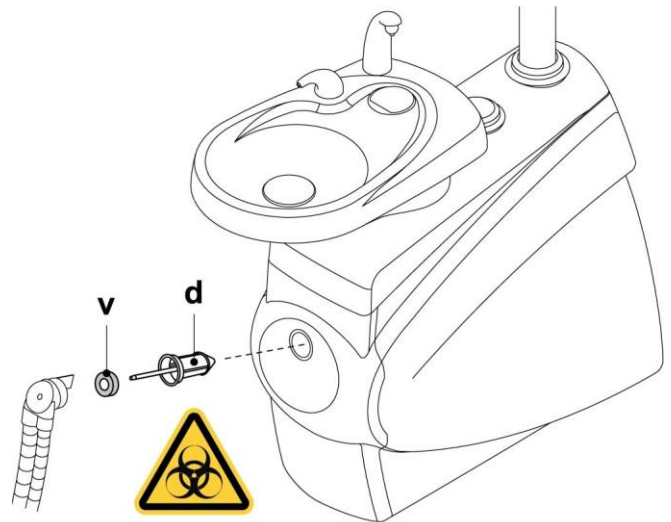
At the end of each work day.

- Suck in 1 litre of water with each suction tube, alternating water and air (keep the suction tube alternately in and out of the water).
- Once rinsed with water, suck in approximately half a litre of the solution prepared with the selected disinfectant with each of the suction tubes used.



NOTE:

after disinfecting, it is good practice to suck in air only to dry the entire suction system (5 minutes).



Every 15 days.

- Clean the separator container and probes with a soft sponge and neutral detergent.
- Clean the drain valve for the separator's container with the device provided for this purpose.

Once a year.

- By technician: check the siphons and drains, check all the internal tubes and plastic and rubber parts subject to wear.

Before leaving the surgery empty for a few days.

- Start suction and run it for 20-30 minutes without sucking in any liquids. The aspirator will dry completely. As a result, salt caused by moisture and basic substances will not form. Said salt may cause fan seizure and motor blockage.

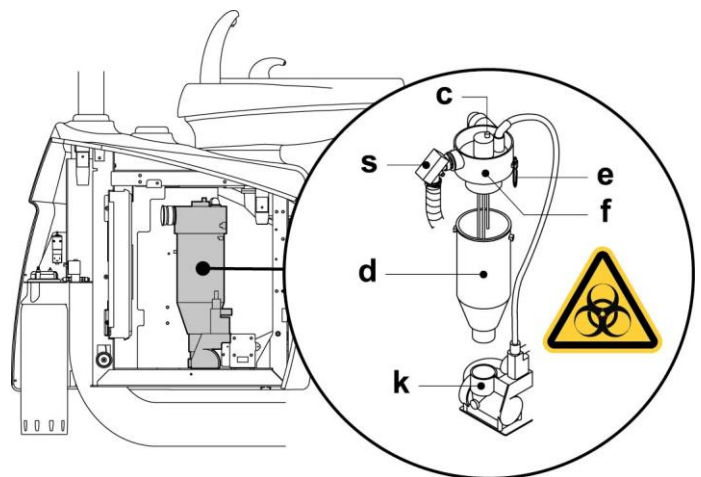
How to remove the separator's container.



WARNING:

Always wear goggles and gloves to prevent contact with infected material when carrying out the following operation.

- Move the dental chair fully up.
- Open the water unit side cover (see paragraph 7.5.).
- Completely empty the separator container, pressing the timed button (c) located on the cover.
- If present, remove the valve (s) for central systems.
- Turn and raise the container (d) until it is detached from the drain pump (k).
- Detach the container (d) from the cover (f) pulling up the two side rubber bands (e).
- After the cleaning operations, refit the container (d) after lubricating the O-rings with **S1-Protective Lubricant for O-Rings** (CEFLA s.c.).
- Lastly, close the water unit side cover.



9.6. CATTANI GRAVITY SURGICAL SEPARATOR

Draining the separator's container.

- Move the dental chair completely up so that as much liquid as possible can be drained from the container.



WARNING:

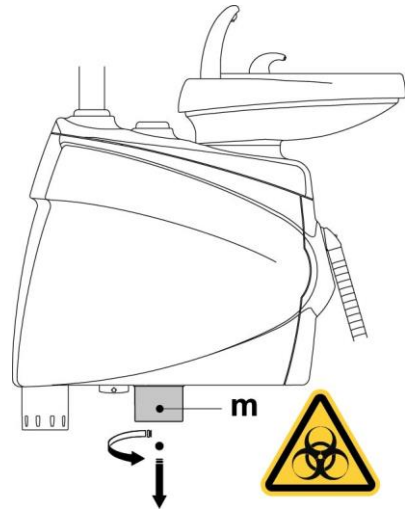
Always wear goggles and gloves to prevent contact with infected material when carrying out this operation.

- Turn the container (**m**) counter-clockwise and remove it.
- Refer to the instructions provided by CATTANI enclosed with the equipment to drain the container in the disposable container provided for this purpose (article code 97290027).



WARNING:

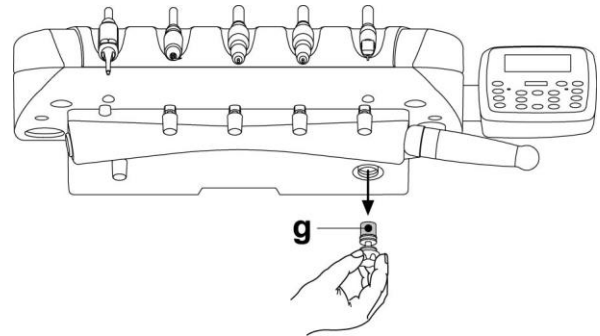
When disposing of disposable containers full of amalgam, observe current local and national laws.



9.7. CLEANING THE TURBINE RETURN AIR FILTER

Monthly check the oil container filter (**g**) present in the turbine's return air line.

If necessary, replace the filter element (code 97290014).



9.8. METASYS AMALGAM SEPARATOR

The maintenance instructions for the METASYS amalgam separator are enclosed with the equipment, if the equipment comes with this type of separator.

The separator control device is located in the water unit.



WARNING:

Always wear goggles and gloves to prevent contact with infected material when cleaning the separator.



WARNING:

When disposing of disposable containers full of amalgam, observe current local and national laws.



9.9. DÜRR AMALGAM SEPARATOR

The maintenance instructions for the DÜRR amalgam separator are enclosed with the equipment if the equipment comes with this type of separator.

The separator control device is located in the water unit.



WARNING:

Always wear goggles and gloves to prevent contact with infected material when cleaning the separator.



WARNING:

When disposing of disposable containers full of amalgam, observe current local and national laws.

9.10. DENTAL CHAIR

The dental chair does not need any special maintenance.

In any case, operation should be checked by CASTELLINI authorised personnel once a year.



10. TECHNICAL DATA

Installation plan:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO	97042090
	PUMA ELI R CART ISO	97042091
	PUMA ELI A R PUMA ELI A R CP	97042105
Technical manual:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	97071175
	PUMA ELI A R PUMA ELI A R CP	97071206
Dental unit spare parts catalogue:		97023126
Dental chair spare parts catalogue:		97023126
Maximum dental unit weight:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	80 Kg.
	PUMA ELI A R PUMA ELI A R CP	90 Kg.
Maximum dental chair weight:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	80 Kg.
	PUMA ELI A R PUMA ELI A R CP	100 Kg.
Maximum dental chair load capacity:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	190 Kg.
	PUMA ELI A R PUMA ELI A R CP	160 Kg.
Rated voltage:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	230V~
	PUMA ELI A R PUMA ELI A R CP	230V~ / 115V~
Nominal frequency:		50/60 Hz.
Absorbed power:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO	1500 W
	PUMA ELI R CART ISO	500 W
	PUMA ELI A R PUMA ELI A R CP	1500W (230V~) 1000W 115V~
Air connection:		1/2 Gas.
Air supply pressure:		6-8 bar.

Air flow rate:	82 l/min.	
Water connection:	1/2 Gas.	
Water supply pressure:	3-5 bar.	
Water delivery rate	10 l/min:	
Water consumption:	2 l/min.	
Water hardness:	< 25 °f (14 °d).	
Drain connection:	ø40 mm.	
Drainage flow rate:	10 l/min.	
Drain duct inclination:	10 mm/m.	
Suction connection:	ø40 mm.	
Suction vacuum (minimum):	65 mbar.	
Suction flow rate:	450 l/min.	
Type-approval:	CE 0051	
Electrical work in compliance with:	IEC 60364-7-710	
Dental unit packaging dimensions:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO	1580 x 780 x 1330(h)
	PUMA ELI R CART ISO	1030 x 680 x 650(h)
	PUMA ELI A R PUMA ELI A R CP	1580 x 780 x 1330(h)
Dental chair packaging dimensions:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	1520 x 730 x 1030(h)
	PUMA ELI A R PUMA ELI A R CP	1520 x 730 x 1030(h)
Dental unit packaging weight:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO	140 Kg.
	PUMA ELI R CART ISO	55 Kg.
	PUMA ELI A R PUMA ELI A R CP	150 Kg.
Dental chair packaging weight:	PUMA ELI R PUMA ELI R CP PUMA ELI R CART PUMA ELI R ORTHO PUMA ELI R CART ISO	150 Kg.
	PUMA ELI A R PUMA ELI A R CP	180 Kg.

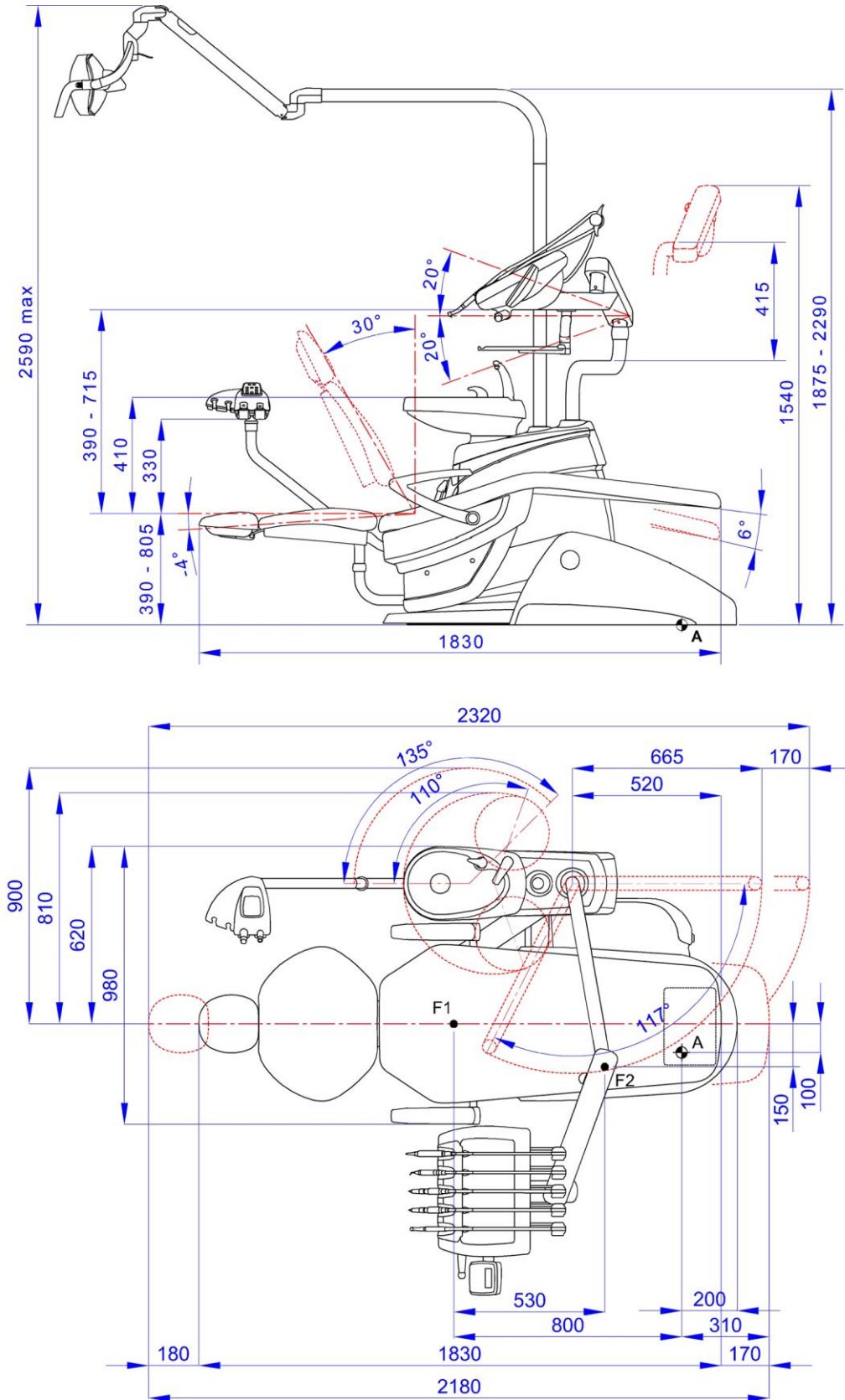
FUSES

Identification	Value	Protection	Position
<i>Dental unit.</i> Fuse F2	T 8 A T 10 A	230 V~: Dental unit power supply line. 115 V~: Dental unit power supply line.	Connection box. Connection box. Connection box. Connection box.
Fuse F4	T 6.3 A	Secondary protection: Water unit.	
Fuse F5	T 6.3 A	Secondary protection: Dental unit.	
Fuse F6	T 6.3 A	Secondary protection: Operating light.	
<i>Dental chair.</i> Fuse F1	T 4 A	230 V~: Dental chair power supply line.	
<i>Quick-couplings.</i> Fuse	T 2 A	230 V~: Power outlet supply line.	Connection box.
<i>MONITOR power supply.</i> Fuse	T 4 A	21 V~ : MULTIMEDIA power line.	Dental chair card area.

* Technical specifications refer to models: PUMA ELI R, PUMA ELI R CP, PUMA ELI R CART, PUMA ELI R ORTHO, PUMA ELI R CART ISO, PUMA ELI A R , PUMA ELI A R CP.

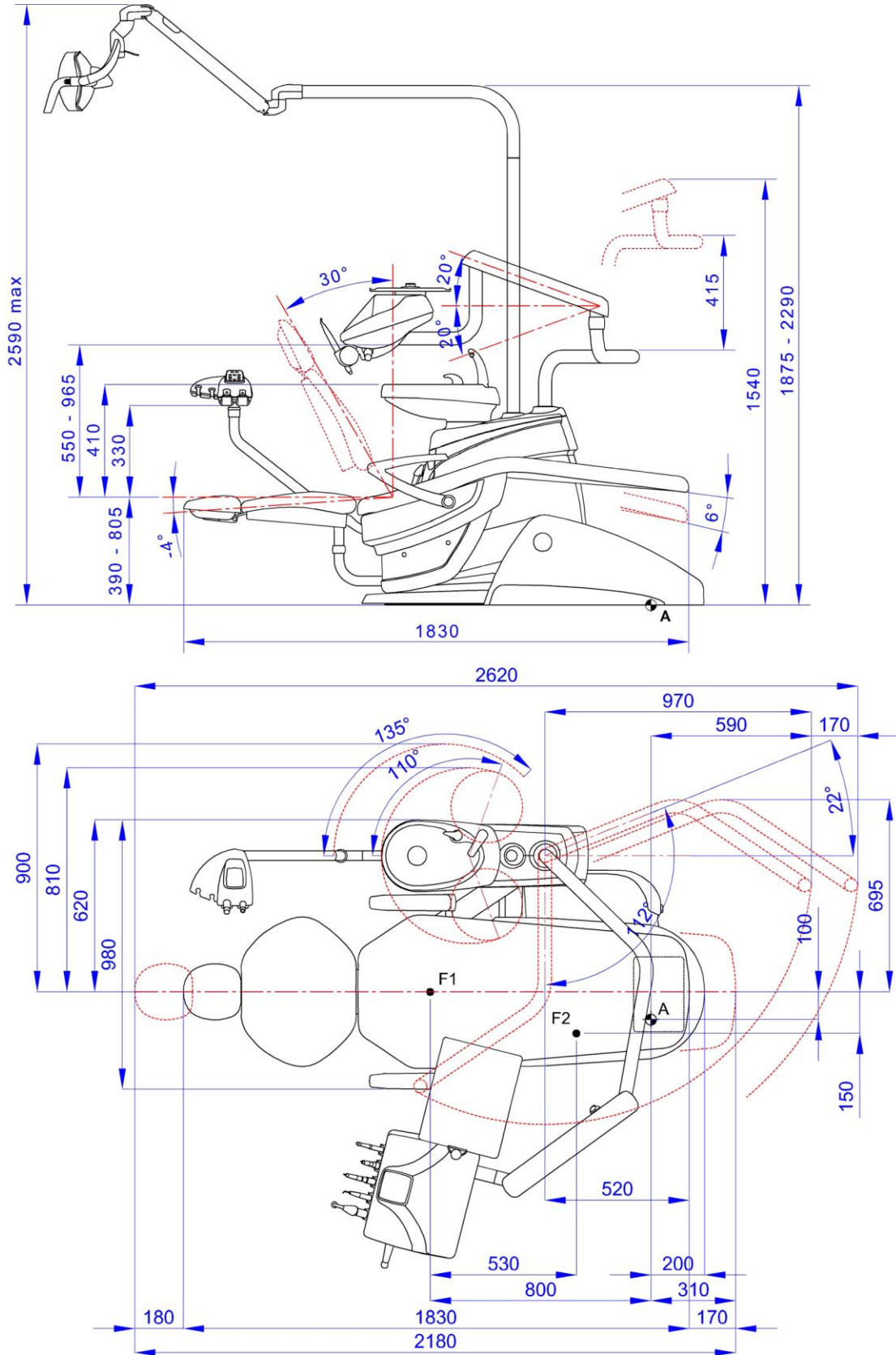
10.1. DIMENSIONAL CHARACTERISTICS

PUMA ELI R DIMENSIONAL CHARACTERISTICS



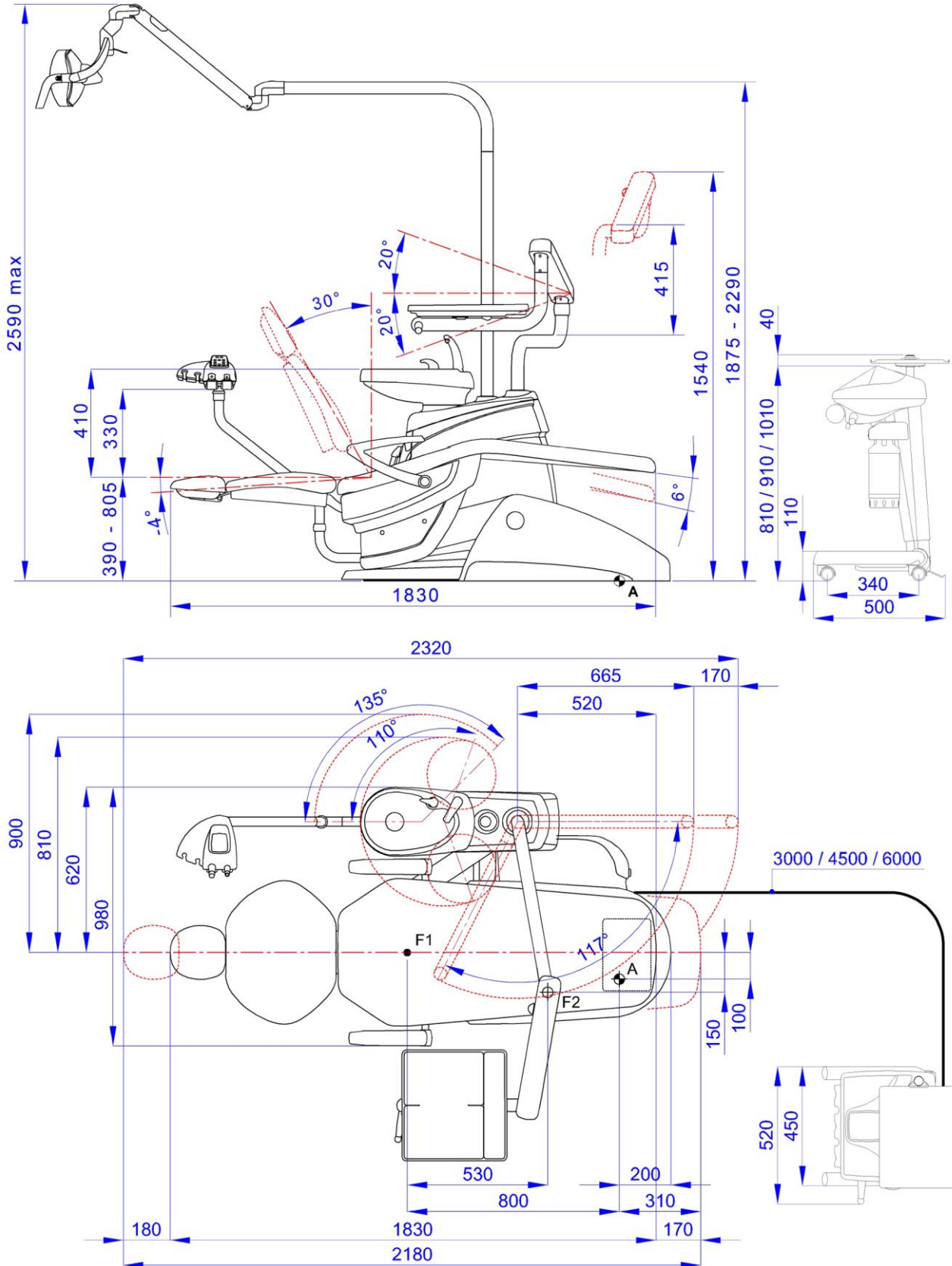


PUMA ELI R CP DIMENSIONAL CHARACTERISTICS



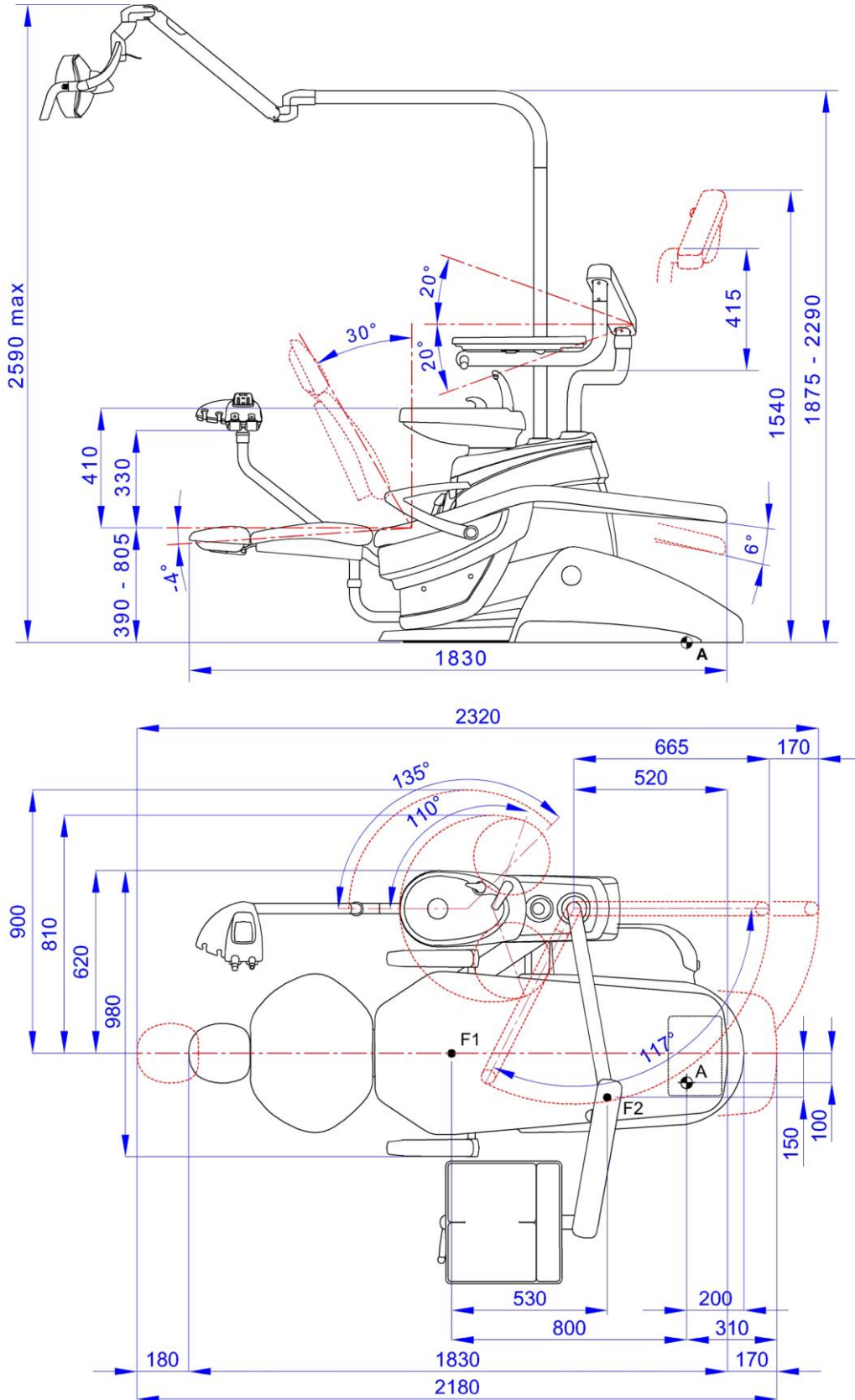


PUMA ELI R CART DIMENSIONAL CHARACTERISTICS



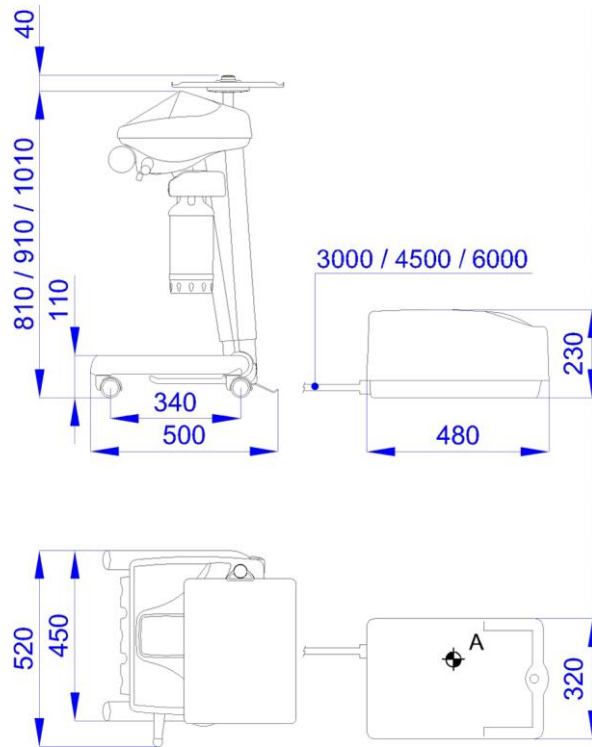


PUMA ELI R ORTHO DIMENSIONAL CHARACTERISTICS



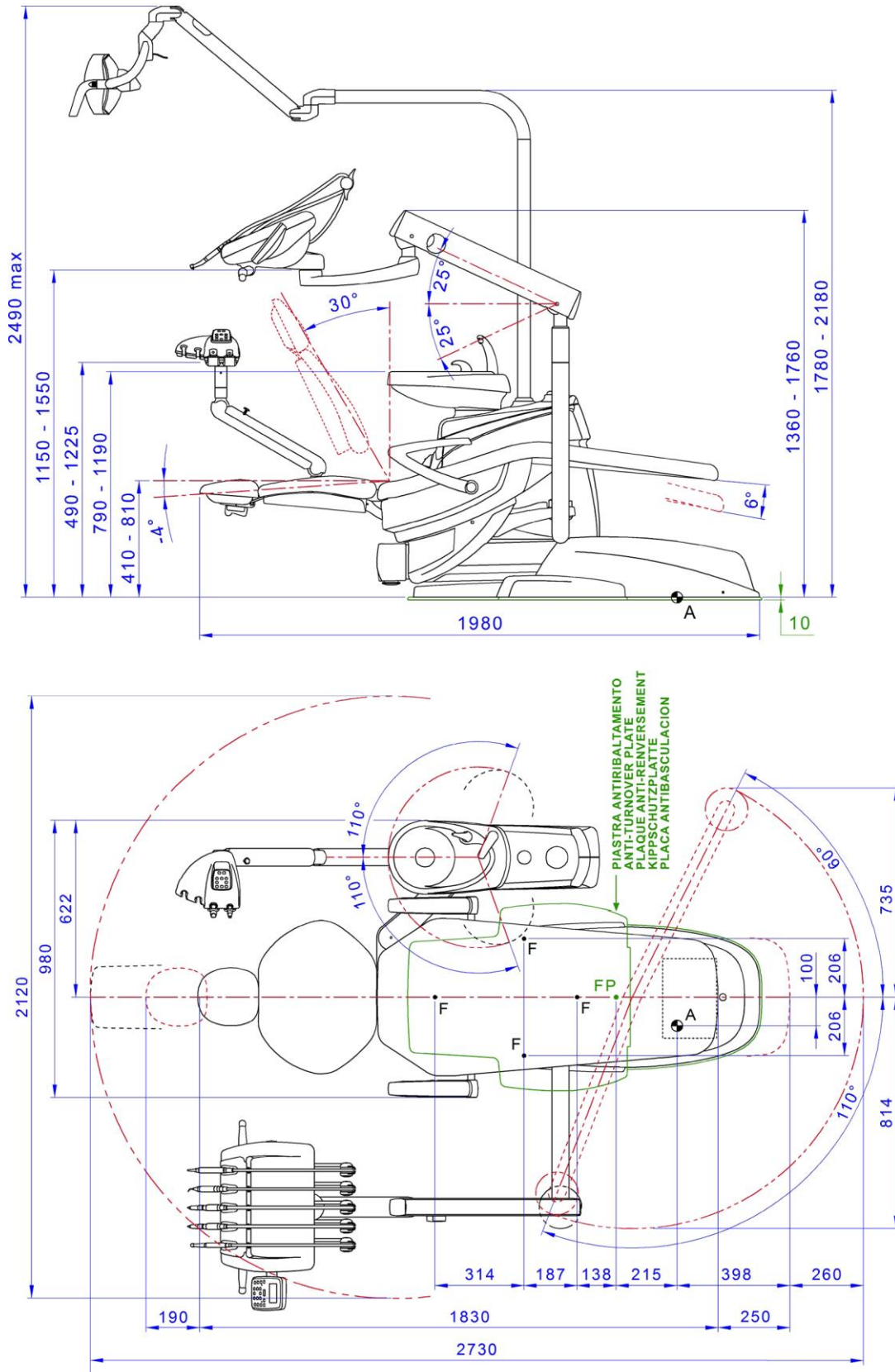


PUMA ELI R CART ISO DIMENSIONAL CHARACTERISTICS



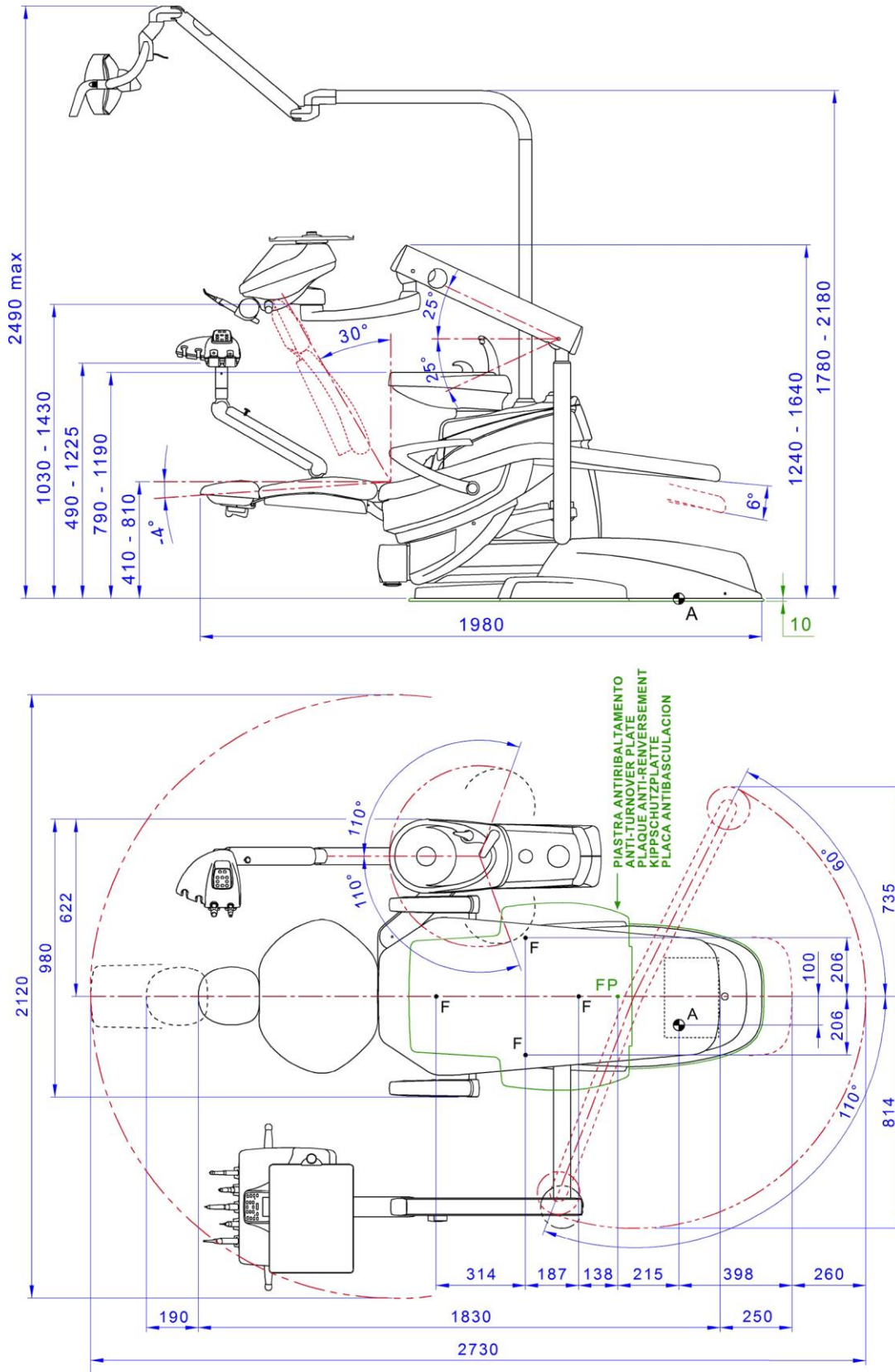


PUMA ELI A R DIMENSIONAL CHARACTERISTICS





PUMA ELI A R CP DIMENSIONAL CHARACTERISTICS





11. DENTAL UNIT OPERATING UNIT MAINTENANCE PLAN

WHEN	PART	ACTION	REFERENCE PARAGRAPH
Daily activity start.	Drain condensate.	Eliminate condensate from air ducts.	See paragraph 9.2.
	Spray ducts (if the dental unit is left off for more than 48 hours).	Disinfection of spray water ducts with manual cycle (if the S.S.S. system is fitted).	See paragraph 7.2.2.
		Disinfection of spray water ducts with semi-automatic cycle (if the AUTOSTERIL system is fitted).	See paragraph 7.3.
	CATTANI suction separator.	Insert inside each suction filter a tablet of VF CONTROL PLUS.	See paragraph 9.5.
	Disposable infection control sheaths.	Apply flexible sheaths and disposable barriers on dental chair and unit.	/
	Instruments.	Lubricate turbine and micromotor handpieces.	See documentation enclosed with instrument.
After each patient.	Spray ducts.	Wash spray water ducts with TIME FLUSHING system or carry out a 20 second manual drainage for each instrument.	See paragraph 7.4.
	Instruments.	Sterilise or disinfect outside.	See relevant paragraphs.
	Surgical suction tubes.	Aspirate about half a litre of 6% diluted STER 3 PLUS solution with each of the cannulas used.	See paragraph 9.4.
	Flexible protections.	Clean, disinfect or sterilise the contaminated flexible protections.	/
	Disposable barriers.	Replace disposable barriers.	/
	Contaminated surfaces.	Clean and disinfect surfaces with STER 1 PLUS .	/
At the end of the work day.	Spray ducts.	Disinfection of spray water ducts with manual cycle (if the S.S.S. system is fitted).	See paragraph 7.2.2.
		Disinfection of spray water ducts with semi-automatic cycle (if the AUTOSTERIL system is fitted).	See paragraph 7.3.
	Bowl filter.	Clean filter in running water. The content must be disposed of separately.	See paragraph 7.1.
	Bowl.	Clean with off-the-shelf detergents formulated for ceramic materials. Do not use acid or harsh products.	See paragraph 7.1.
	Surgical suction filter.	Clean or replace filter.	See paragraph 9.3.
	Surgical suction tubes.	Aspirate about half a litre of 6% diluted STER 3 PLUS solution with each of the cannulas used.	See paragraph 9.4.
	Hydraulic saliva ejector.	Clean the filter of the saliva ejector terminal.	See paragraph 6.6.
	Flexible protections.	Clean, disinfect or sterilise the flexible protections.	/
	Disposable barriers.	Eliminate all used disposable barriers.	/
	Dental unit and chair surfaces.	Clean and disinfect surfaces with STER 1 PLUS .	/
	When needed.	CATTANI suction separator.	Clean the separator container, drain valve and probes.
METASYS amalgam separator.		Empty or replace separator container.	See documentation enclosed with separator.
DÜRR amalgam separator.		Empty or replace separator container.	See documentation enclosed with separator.
Operating light. VENUS, VENUS PLUS		Clean transparent screen, reflector and handles.	Refer to the CD-ROM user's manual.
VENUS PLUS-L operating lamp.		Clean transparent screen and handles.	See paragraph 8.1.1.
VENUS e operating lamp.		Clean transparent screen and handles.	See paragraph 8.1.2.
Monitor on light pole.		Clean surfaces.	See documentation enclosed with monitor.
Coated surfaces and upholstery.		Clean and disinfect surfaces with STER 1 PLUS .	See paragraph 1.4.
Weekly.	Suction tube holder terminals.	Lubricate the O-ring.	See paragraph 9.4.
Monthly.	Turbine return air filter.	Check the filter and replace it if necessary (code 97290014).	See paragraph 9.7.
Yearly.	Dental chair.	Contact the technical service department for general inspection.	/

**DICHIARAZIONE DI CONFORMITÀ “CE / EU” / “CE / EU” CONFORMITY DECLARATION
DECLARATION DE CONFORMITÉ “CE / EU” / ERKLÄRUNG VON “CE / EU” ZUSTIMMUNG / DECLARACION DE CONFORMIDAD “CE / EU”
DECLARAÇÃO DE CONFORMIDADE “CE / EU” / ΔΗΛΩΣΗ ΠΙΣΤΟΤΗΤΑΣ “CE / EU” / ДЕКЛАРАЦИЯ СООТВЕТСТВИЯ “CE / EU”
DEKLARACJA ZGODNOSCI WE “CE / EU” / “CE / EU” UYGUNLUK BELGESİ**

Prodotto tipo/ Product type :

Matr./ Serial N°:

Incollare in questo spazio l'etichetta del complesso odontoiatrico o di altra apparecchiatura o indicare modello e numero di matricola
Stick the label of the dental equipment or other device into this space or write model and serial number

- I** Dichiaro sotto la nostra esclusiva responsabilità che i prodotti ai quali questa dichiarazione si riferisce sono conformi
1) ai requisiti essenziali (Allegato I) presenti nella direttiva 93/42/CEE Dispositivi Medici (D.Lgs.46/97) e successive modifiche ed integrazioni (**dispositivo medico di Classe IIa**)
2) alla direttiva 2011/65/UE del Parlamento europeo e del Consiglio dell'8 giugno 2011, sulla restrizione dell'uso di determinate sostanze pericolose nelle apparecchiature elettriche ed elettroniche (Rohs 2)
- GB** We declare, on our sole responsibility, that the products referred to herein are in compliance with
1) the essential requirements (Annexe I) of Directive 93/42/EEC Medical devices (Leg. Decree 46/97) and subsequent amendments and integrations (**Class IIa medical device**)
2) Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (Rohs 2)
- F** Nous déclarons, sous notre complète responsabilité, que les produits auxquels la présente déclaration fait référence sont conformes
1) aux exigences essentielles (Annexe I) présentes dans la directive 93/42/CEE "Dispositifs médicaux" (Décr.L. 46/97) et modifications successives et intégrations (**dispositif médical de Classe IIa**)
2) à la directive 2011/65/UE du Parlement européen et du Conseil du 8 juin 2011 relative à la limitation de l'utilisation de certaines substances dangereuses dans les équipements électriques et électroniques (Rohs 2)
- D** Wir erklären hiermit in alleiniger Verantwortung, dass die Produkte, auf die sich diese Erklärung bezieht, konform sind mit
1) den grundlegenden Anforderungen (Anhang I) der Richtlinie 93/42/EWG über Medizinprodukte (Gesetzesverordnung 46/97) und nachfolgenden Änderungen und Ergänzungen (**medizinisches Gerät der Klasse IIa**)
2) der Richtlinie 2011/65/EU des Europäischen Parlaments und des Rats vom 8. Juni 2011 zur Beschränkung der Verwendung bestimmter gefährlicher Stoffe in Elektro- und Elektronikgeräten (Rohs 2)
- E** Declaramos bajo nuestra exclusiva responsabilidad que los productos a los que esta declaración se refiere, están conformes con
1) los requisitos esenciales (Anexo I) presentes en la directiva 93/42/CEE Dispositivos Médicos (D. Leg. 46/97) y sucesivas modificaciones e integraciones (**dispositivo médico de Clase IIa**)
2) la directiva 2011/65/UE del Parlamento europeo y del Consejo del día 8 de junio de 2011, sobre la restricción del uso de determinadas sustancias peligrosas en los aparatos eléctricos y electrónicos (Rohs 2)
- P** Declaramos sob a nossa exclusiva responsabilidade que os produtos aos quais esta declaração se refere estão em conformidade
1) com os requisitos essenciais (Anexo I) presentes na directiva 93/42/CEE Dispositivos Médicos (em Itália, transposta pelo Decreto Legislativo 46/97) e posteriores alterações e aditamentos (**dispositivo médico de Classe IIa**)
2) com a directiva 2011/65/UE do Parlamento europeu e do Conselho de 8 de junho de 2011, relativa à restrição do uso de determinadas substâncias perigosas em equipamentos elétricos e eletrónicos (Rohs 2)
- GR** Δηλώνουμε με την αποκλειστική ευθύνη μας ότι τα προϊόντα στα οποία αναφέρεται η παρούσα δήλωση είναι σύμφωνα
1) με τις βασικές απαιτήσεις (Προσάρτημα I) της οδηγίας 93/42/ΕΟΚ Ιατροτεχνολογικών Προϊόντων (Ν. Διάτ.46/97) και μεταγενέστερες τροποποιήσεις και συμπληρώσεις (**ιατροτεχνολογικό προϊόν Κατηγορίας IIa**)
2) με την οδηγία 2011/65/ΕΕ του Ευρωπαϊκού Συμβουλίου και του Συμβουλίου της 8 Ιουνίου 2011, για τον περιορισμό της χρήσης ορισμένων επικίνδυνων ουσιών σε ηλεκτρικό και ηλεκτρονικό εξοπλισμό (Rohs 2)
- PY** Под нашу исключительную ответственность заявляем, что изделия, к которым относится данная декларация, соответствуют
1) основным требованиям (Приложение I) директивы 93/42/ЕЭС Медицинские устройства (Законодательный указ № 46/97) и последующим изменениям и дополнениям (**медицинское устройство Класса IIa**)
2) директиве 2011/65/ЕС Европарламента и Совета Европы от 8 июня 2011 года по ограничению использования определенных опасных веществ в электрическом и электронном оборудовании (Rohs 2)
- PL** Oświadczamy na swoją wyłączną odpowiedzialność, że produkty objęte niniejszym oświadczeniem są zgodne:
1) z zasadniczymi wymaganiami (Załącznik I) przewidzianymi dyrektywą 93/42/EWG Wyroby Medyczne (D. z mocą ustawy 46/97) wraz z późniejszymi zmianami i uzupełnieniami (**wyrób medyczny Klasa IIa**)
2) z dyrektywą 2011/65/WE Parlamentu europejskiego i Rady z dnia 8 czerwca 2011r. w sprawie ograniczeń we wprowadzaniu do obrotu i stosowaniu w sprzęcie elektrycznym i elektronicznym określonych niebezpiecznych substancji (Rohs 2)
- TR** Bu beyannameye bahsi geçen ürünlerin aşağıda belirtilenlere uygun olduğunu kendi mühasır sorumluluğumuz altında beyan ederiz:
1) (Kanun hükmünde Kararname 46/97) Medikal Aygıtlar 93/42/CEE direktifinde mevcut (Ek 1) ana gereklilikler ve sonraki değişiklikler ve eklemelerde belirtilenler (**IIa sınıf medikal aygıt**)
2) 8 Haziran 2011 tarihli Avrupa Parlamentosu ve Konseyi'nin "Elektrikli ve elektronik cihazlarda bazı tehlikeli maddelerin kullanılmasına ilişkin kısıtlamalar" 2011/65/UE direktifi (Rohs 2)

Imola, li _____

Bussolari Paolo
Managing Director

