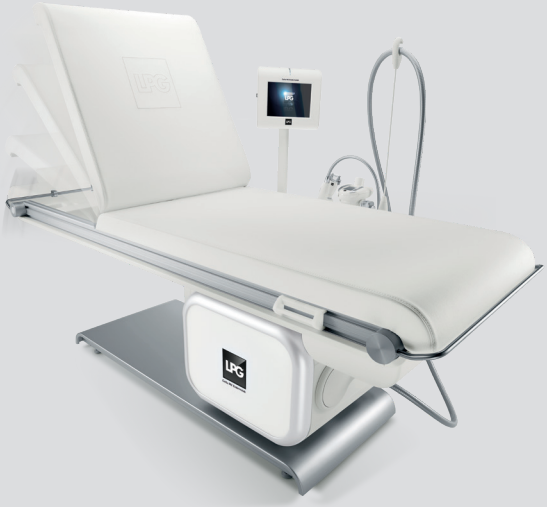


Cellu M6® Endermolab

OPERATOR'S MANUAL

Please read the complete manual carefully before using your equipment

GU 0904 - EN
Edition D dated 10/2020

Congratulations on the purchase of your device Cellu M6® Endermolab i device.
This model represents many years of research in the design and production of cutaneous tissue treatment systems. You will be able to fully appreciate the technical perfection and reliability that have made LPG Systems the leader in this field. This operator's manual contains the operating description, basic maintenance instructions to be performed periodically and the safety instructions.
Your device is intended for use in the treatment of connective tissue. It should be used only by a professional who has attended the manufacturer's training provided by LPG Systems or an approved provider if you live outside of France.
If you have any doubts whatsoever concerning the operation or maintenance of your equipment, please contact LPG Systems via your distributor.

ATTENTION

In order to respond better to customer requirements and expectations, LPG Systems is continuously researching ways of improving the design and quality of its products. This will explain the few minor differences between your equipment and the item described in this guide.

→ PACKAGE CONTENTS

- One Cellu M6® Endermolab *i* device
- One main Ergodrive™ head
- One Keymodule™ set (KM80)
- One TR50 head
- One set of auxiliary heads (TR15 et TR30)
- One set of micro-nozzles/micro-heads
- One Ergolift head
- Two Ergolift™ chambers (Lift 20 and Lift 10)
- One set of Lift heads
- One operator's manual
- Two electrical power cords
- One POS marketing set

Depending on the version you have (see serial number on the nameplate; "B", "2i" or "i"), some protocols are not activated and their accessories are not provided. Accordingly, the paragraphs describing them do not concern this version (see table on the next page).

↗ ATTENTION

Unpacking instructions to be retained.

	Version 2 i	Version i	Version i B
Ergodrive	√	√	√
KM80	√	√	√
TR50	√	√	√
TR30, TR15	√	√	√
Micro-nozzles Micro-heads	√	√	√
Ergolift Lift 20 Lift 10	√		
Lift heads		√	
TML30 TML20 TML10		√	
GU	√	√	√
Power cords	√	√	√
POS	√	√	√

→ TABLE OF CONTENTS

→ CELLU M6® ENDERMOLAB

INTENTED USE

The CELLU M6® Endermolab i Medical device is intended for therapeutic use.
It can be used to:

1. Reduce secondary lymphedema of the arm after a mastectomy
2. Improve secondary lymphedema
3. Improve lymphatic circulation in the treated area
4. Relieve minor muscle aches and pains
5. Relieve muscle spasms
6. Improve local blood circulation
7. Temporarily relieveof the minor muscular pain associated with DOMS (Delayed Onset Muscle Soreness)
8. Improve local circulation during burn rehabilitation
9. Reduce the appearance of cellulite and the circumference of treated areas
10. Temporarily improve lymphatic circulation and local blood circulation to improve skin trophicity in the treated areas
11. Improve skin quality, scars and fibrosis
12. Improve skin aging (wrinkles, fine lines, skin sagging, fat infiltration, firmness, elasticity, complexion and eye bags)
13. Stimulate fibroblasts (collagen, elastin and hyaluronic acid synthesis)

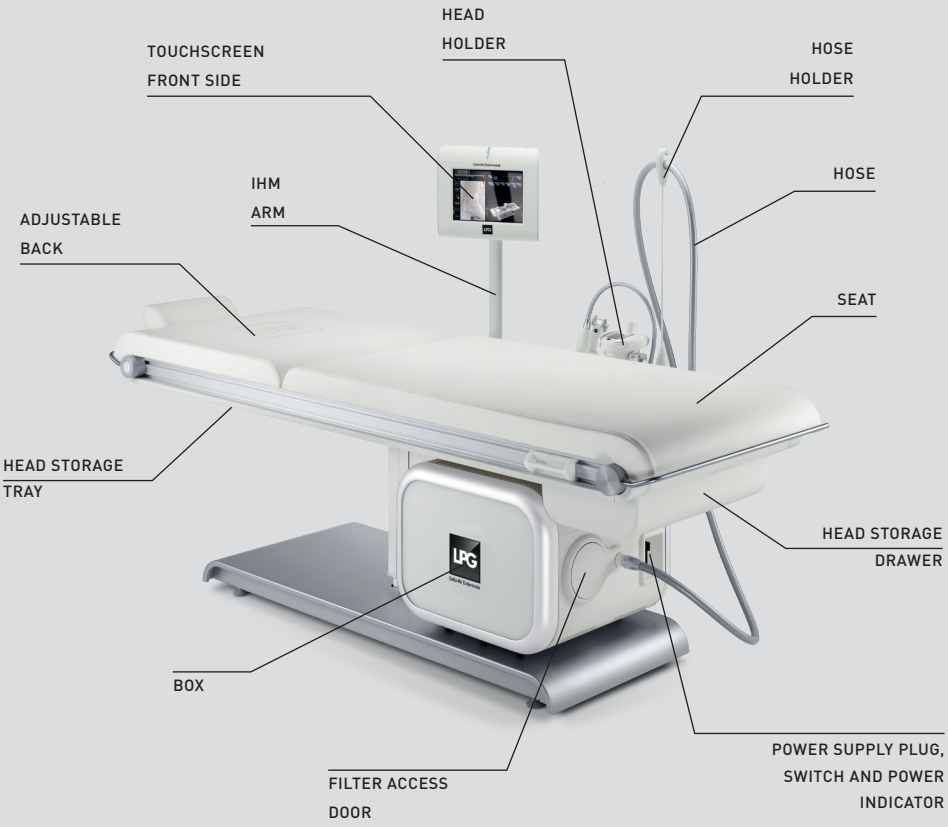
It utilizes mecano-stimulation treatment heads for bodycontouring and anti-aging applications. The device can be used in hospitals, therapy centers and institutions by specialists and physiotherapists. It is an independent device that cannot be combined with other devices. It is to be used by professionals who are specially trained by LPG Systems in the use of the device and is not suitable for home use. The device should be used on adults only.

OPERATING PRINCIPLES

The operating principles of the Cellu® M6 Endermolab i medical device consist of a suction force coupled with movements of rolls/valves, performed with treatment heads.

These heads are placed on the healthy skin of the patient and then moved across the area to be treated by the professional who has been trained by LPG Systems.

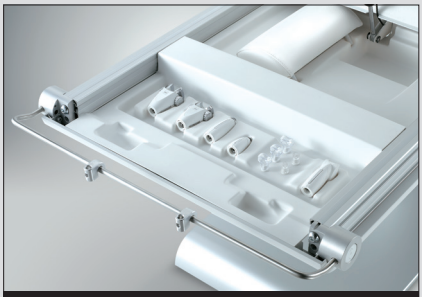
→ CELLU M6® ENDERMOLAB (CONT'D)



ATTENTION

The device can only operate if it is connected to the power supply by its power cord and provided the ON switch has been actuated and the green voltage light is on.
After switching on the unit, please wait a few seconds for the screen to display information.

→ HEAD STORAGE TRAY AND DRAWER



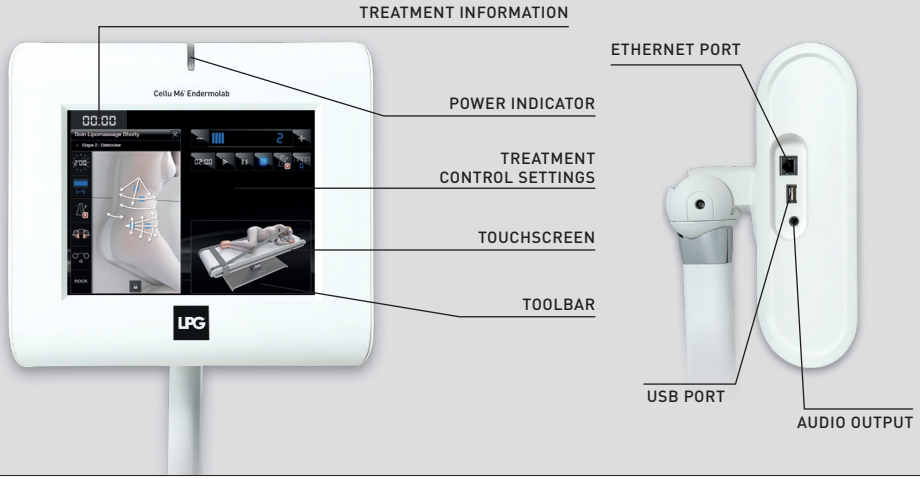
HEAD STORAGE TRAY

HEAD STORAGE DRAWER

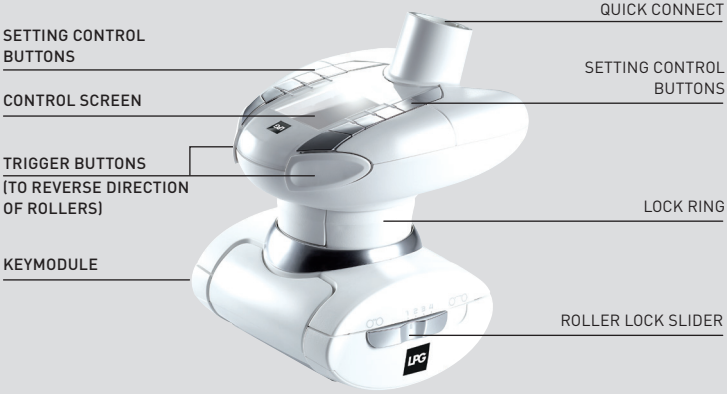
→ FILTER ACCESS

THE FILTER IS ACCESSIBLE ON THE SIDE OF THE BOX.

→ CONTROL SCREEN



→ ERGODRIVE™ HEAD

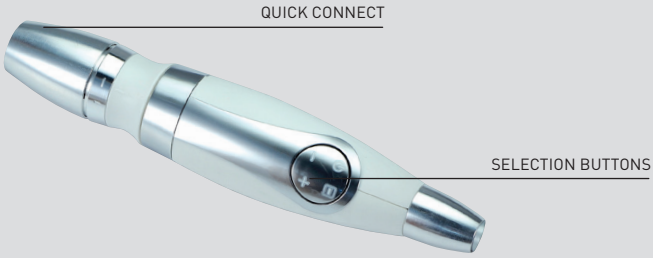


↗ ATTENTION

For detailed instructions on using the touch interface, refer to the touch interface operating manual received during training and available from customer service.

→ TR50 HEAD

→ ADAPTER



→ HEAD

→ CONNECTION BETWEEN BOX AND TABLE

→ ADJUSTABLE BACK

To raise or close the table back, pull the handle on the adjustable back and then raise or lower the back while holding onto the handle.

→ HEIGHT ADJUSTMENT

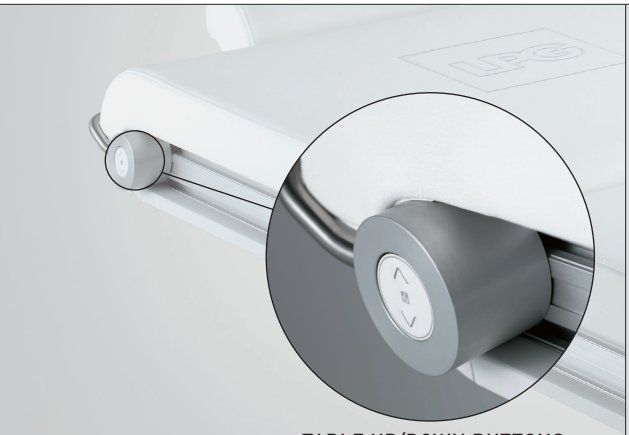


TABLE UP/DOWN BUTTONS

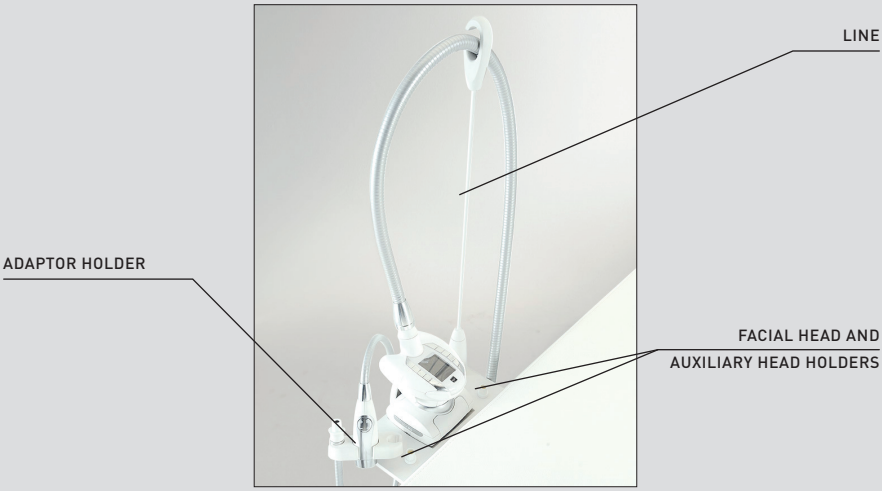
The table height is adjustable using the controls on the table (see photo to left).
Simply press the appropriate buttons to move the column up or down.

→ FRONT SIDE ADJUSTMENT

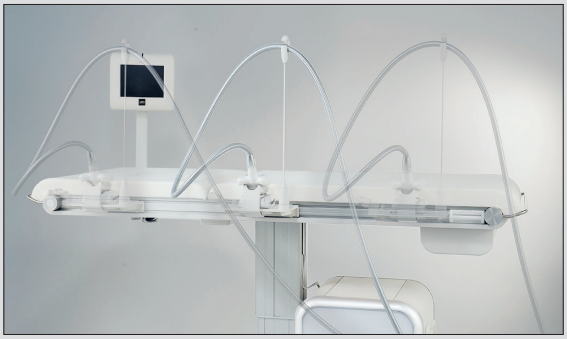
The front side can be adjusted using the articulated dial.
To adjust the front side, simply move it to the desired position.

→ SHELF

The Cellu M6® Endermolab *i* device is equipped with a shelf to hold the different treatment heads



This shelf can be moved along the length of the table by sliding it along the provided rail.



→ STRAPS

Your Cellu M6® Endermolab i device comes with a set of straps, including one lateral strap and two longitudinal straps.
These straps should be set up as shown in the photos below.

Attach the lateral strap:

Attach the longitudinal straps:

↗ ATTENTION

Before use, make sure that the corners are properly secured.

→ IMPORTANT SAFETY INFORMATION

All safety precautions must be observed while using electrical equipment. Please read all safety notices and precautions prior to use of the device.

DANGER - TO MINIMISE THE RISK OF ELECTRICAL SHOCK:

- Always disconnect the device from the electrical supply outlet after use and before cleaning and maintenance.
- Check that the supply voltage of the unit indicated on the data plate complies with the power supply voltage.
- The device must be connected by the power cord[1] supplied to a grounded outlet in accordance with current electrical standards. Electrical adapters must not be used with this device.

→ WARNING

TO MINIMISE THE RISK OF BURNS, FIRE, ELECTRICAL SHOCK OR INJURY

- The device must not be left unattended while connected to the electrical supply.
- Disconnect the device from the electrical supply if it is not going to be used for a long period.
- Special attention is required while using the device with, or in the proximity of, children or disabled persons.
- Never use the device for purposes other than those recommended by LPG Systems. Only use the treatment heads supplied with your device or those recommended by LPG Systems.
- Never use the device if:
  - the electrical power cord or outlet is damaged.
  - the device does not function correctly.
  - the device is damaged or has fallen or been dropped.
  - the device has been exposed to excessive humidity.

  In such cases, return the device to an approved LPG service centre.
- Do not move the unit by pulling the electrical power cord.
- Fully unwind the supply electrical from its support and keep it away from warm surfaces.
- Never use the device if the ventilation ports are obstructed. Ensure that the ventilation ports are kept clear of dust or other contaminants.
- Do not allow solid debris, liquid or other foreign bodies to fall or be sucked into the unit, as these could cause damage.
- Never use the device on a dusty, uneven floor or in a moist atmosphere.
- Never use the device in the presence of aerosols or oxygen.
- Before disconnecting the unit from the electrical supply, set all controls to the 'off' position and unplug the unit.
- It is prohibited to modify this equipment without prior authorisation from LPG Systems.

ATTENTION

[1] **Europe** VII-H05VVF3G1,50-C19 & VII-H05VVF3G1,00-C13; **Italy** I/3/16-H05VVF3G1,50-C19 & I3G-H05VVF3G1,00-C13; **Switzerland** 23G-H05VVF3G1.50-C19 &12G-H05VVF3G1.00-C13 ; **UK** BS13/13-H05VVF3G1,50-C19 & BS13/13-H05VVF3G1,00-C13 ; **Japan** 498GJ-VCTF3X2.00-C19 & 498GJ-VCTF3X1.25-13V ; **USA, Canada, Mexico** N5/15-SJT3X14AWG-C19 & 498G-SJT3X18AWG-C13 (connect to Hospital grade receptacle in hospital environment)

→ SAFETY INFORMATION

ATTENTION: KEEP THESE INSTRUCTIONS

Your device should be used on clean and healthy skin. It is important to read and respect the following precautions and contraindications before using your device.

- Never touch the patient and the device's unprotected cables or connectors simultaneously.
- Never use the auxiliary adapter as a treatment head or allow it to come into direct contact with the skin.
- Do not use the main head to carry out treatment on the scalp.
- Only use treatment heads supplied with your device or recommended by LPG Systems.
- Only use LPG Systems' treatment garments.
- LPG Systems will not be liable for any inappropriate use of the device.
- The operator must be particularly attentive to the sensations felt by the person undergoing treatment.
- The operator must ensure that the parameters (intensity, sequentiality, differential, etc.) are always adapted to the cutaneous tissue being treated.
- Do not put more than 135 kg on the table.
- The controls used to adjust the height of the table can be inhibited if the cable is disconnected.
- When the table is being used without supervision, it should be put in the lowest position to avoid the risk of falling.
- After use, put the table in the lowest position to avoid the risk of falling.

  Do not use the USB and ethernet connections during treatment.
- Do not operate the unit in unsuitable environmental conditions (see technical specifications).
- Do not use the treatment heads directly on the skin. Wear the ENDERMOWEAR™ treatment suit provided by LPG Systems.
- **Service precaution:** risk of explosion if the Microbattery Button Cell located on the HMI board (type 3V 190 mAH, VARTA 6 032 101 501) is replaced by an incorrect type. Dispose of used Cells according to country regulation.

↗ ATTENTION

Any serious incident occurring with your device should be reported to your local LPG distributor and competent authority.

→ CONTRAINDICATIONS

- Do not treat open wounds, eyes, intracavity areas, mucous membranes, genitals or nipples.
- This device is not recommended for pregnant women. In the event of pregnancy, do not treat the lumbar-abdominal region. Consult with a doctor regarding this treatment.
- Do not treat a patient with an infectious disease, a growing tumour, a phlebitis, a wound or an infected area.
- Do not treat a patient with skin cancer, a visible tumour or other cancerous lesions. Consult with a doctor in cases where the patient has a case history of tumours or is in remission.
- Do not treat inflamed or swollen areas or scars from a recent surgery without medical advice and LPG® device training for the affected areas.
- Do not treat patients with circulatory problems without first consulting their doctor.
- Do not treat any swollen or inflamed areas without seeking medical advice and without having had training in specific LPG® device in this particular area.
- Stop treatment immediately if the patient experiences pain and consult a doctor.
- This device should not be used on skin rashes, herpes, inflamed or infected acne or vitiligo.
- To avoid bruising, exercise caution when determining a patient's level of sensitivity and avoid use on patients taking anti-coagulant drugs.
- For a more detailed list of the indications and contraindications of ENDERMOLOGIE, please refer to the training manuals.
- As this list is not exhaustive, always seek out medical advice in the event of doubt.
- Because of the risk of interference, it is important that the professional ensures the patient is not equipped with a personal medical device such as a pacemaker. If this is the case, the professional should obtain details about the device in question to ensure that any interference will not affect the correct use of the equipment.
- Do not use this device on unhealthy/ damaged skin. Avoid affected areas and any area of the body that have undergone plastic surgery until eccymosis, oedema and pain disappear.

ATTENTION

This device contains programs to help the operator obtain the best anticipated results for each case undergoing treatment. Under no circumstances may these programs be construed as a guarantee of successful treatment, which varies depending on the morphology, physiology and eating behaviour of each patient.

→ INDEX

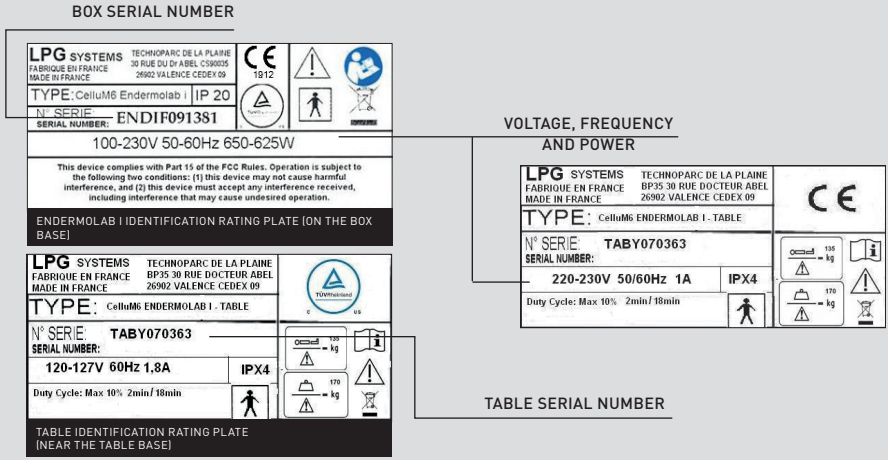
→ IDENTIFICATION RATING PLATE

Your unit is identified by two serial numbers shown on the rating plates.
These rating plates also show the permitted supply voltage for the unit.

If you need to contact LPG Systems because of a technical problem, please indicate the serial numbers of your Cellu M6® Endermolab i device.

These serial numbers provide information on the year and month of manufacture of your unit.

The letter indicates the year the device was manufactured. Z=2009, A=2010, B=2011, etc.
The two digits indicate the production month: 01=January; 02=February; 03=March; etc.



This icon indicates that the unit was sold after August 13, 2006. In conformity with the 2002/96/CE directive, it cannot be thrown away with standard household waste but must be disposed of by means of recycling. When your device reaches the end of its useful life, it must be brought to an appropriate recycling center or returned to your dealer. By doing so, you help the environment by contributing to the conservation of natural resources and the protection of human health.

This icon indicates that some specific warnings or precautions associated with this device are not on the label.

This symbol means always consult the accompagnying documents before using your device.

This symbol indicates the name and address of the manufacturer.

↗ ATTENTION

Identification rating plates may vary. The approved one is one on your machine.


This symbol means that your device has type BF applied parts which come into contact with the patient. These parts are electrically isolated from all of the device's other parts. These applied parts are the treatment heads.


This symbol means store the device somewhere protected from inclement weather.


This symbol indicates temperature limits.


This symbol indicates relative humidity limits.


This symbol means "Danger: High Voltage."


This symbol means "Use under prescription."


This symbol means that the maximum exterior mechanical load permitted on this equipment is 170 kg.


This symbol means that the the maximum patient weight permitted on this equipment is 135 kg.


This symbol means always consult the accompanying documents before using your device.

→ CLEANING THE DEVICE

It is recommended that you clean your unit as often as possible, not onlyfor hygienic and aesthetic reasons, but also because cleaning the unit will help keep it in a good state and extend its useful life.

The device should be cleaned with a damp non-abrasive sponge after each use.

Using a vacuum cleaner with a fine nozzle, clean the following sections:

- Interior of the head storage drawer.
- Interior of the head storage tray.
- Interior of the filter access door.

Using a moist sponge, clean the following sections:

- All the external hoods.
- Hoses.
- The electrical power cord.

Using a cloth soaked with a small amount of alcohol-free domestic cleaning product, clean the following sections:

- Control screen and control panel.
- Interior of the head storage drawer.
- Interior of the head storage tray.
- Interior of the filter access door.

↗ ATTENTION

Do not use corrosive products such as acetone, trichloroethylene or rubbing alcohol.

→ REPLACING THE FILTER CARTRIDGE AND FOAM

Your device contains one filter cartridge and one filter foam.
These components guarantee the efficiency of your unit and prolong its useful life.

Ensure that the filter cartridge is changed as soon as the command screen displays one of these messages **(Fig. 1-2)**.

Icon indicating a filter change is required **(Fig. 2).**

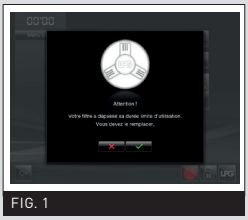


FIG. 1

FIG. 2

CHANGE THE FILTER

Access the 'filter change' menu as follows:

Select the 'maintenance' menu by pressing the icon indicated **(Fig. 3)**.

Select the 'filter' menu by pressing the icon indicated **(Fig. 4)**.

FIG. 3

PRESS THIS ICON

FIG. 4

PRESS THIS ICON

The 'filter change' screen indicates which filter requires changing : **(Fig. 5)**.

Once the filter cartridge is replaced, the filter meter should be reset by pressing the icon indicated : **(Fig. 6)**.

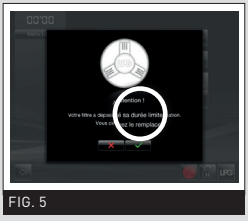


FIG. 5

FILTER LIMIT EXCEEDED

FIG. 6

FILTER METER

→ REPLACING THE FILTER CARTRIDGE

Your device contains one filter cartridge to guarantee the efficiency of your device and to prolong its life.

Be sure to change it as soon as the screen displays a warning message.

Open the filter access door. Unscrew the filter cartridge and replace it with a new one.

Contact LPG Systems Customer Support to stock up on filter cartridges so that you always have spares.



→ REPLACING THE FILTER FOAM

Your device contains two filter foams to guarantee the efficiency of your device and to prolong its life.

Be sure to change them as soon as the screen displays a warning message.

1. Move the cover downward by sliding your hands into the designated notches (as shown in the photo below).
2. Unclip the cover.
3. Replace the filter foams with new ones.

↗ ATTENTION

The device must never be used without a filter. If no filter is fitted, the device must be switched off.

→ CONNECTING/DISCONNECTING THE MOTORIZED TREATMENT HEADS

To **connect** the heads to the hose, follow the procedure below:

Position the locking ring in the locked position **(Fig. 1)**.

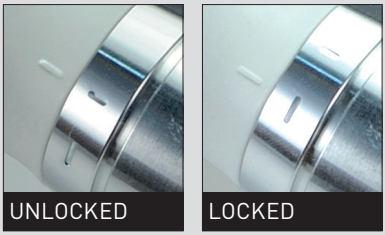
Position the end of the hose so that the hose key is lined-up with the key slot of the treatment head connection **(Fig. 2)**.

Push the hose into the treatment head connection until it clicks into place.

To **disconnect** the heads, position the locking ring in the unlocked position **(Fig. 3)**.

Pull the locking ring towards the hose **(Fig. 4)**.

Carefully remove the hose by pulling it by the white ring **(Fig. 5)**.



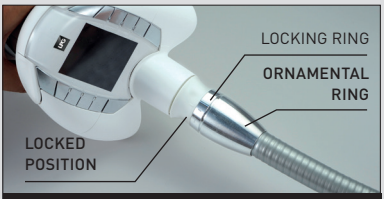


FIG. 1

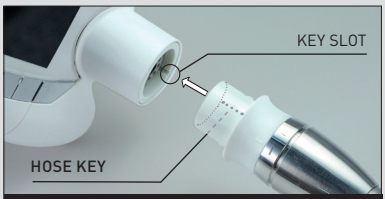


FIG. 2

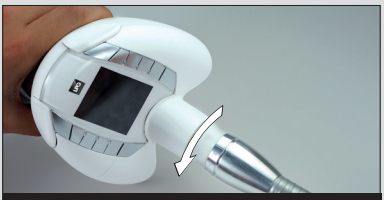


FIG. 3

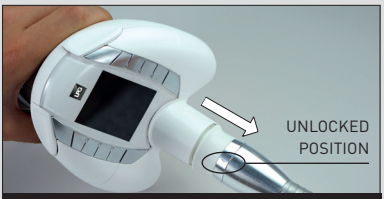


FIG. 4

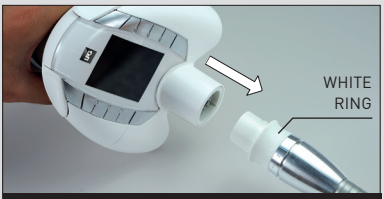


FIG. 5

→ INSTRUCTIONS FOR DISCONNECTING THE HOSE

The hose can be disconnected from the box simply by following the procedure described below:

- Unlock the connection by turning the aluminum ring **(Fig. 1)**.
- Lift the aluminum ring **(Fig. 2)**.
- Gently remove the hose by pulling on the white ring **(Fig. 3)**.

FIG.1

FIG.2

FIG.3

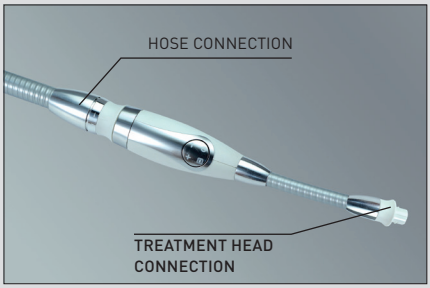
HOSE DISCONNECTED

→ CONNECTING/DISCONNECTING THE ADAPTER

To connect or disconnect the hose adapter, follow the procedures below.

Only the auxiliary heads, the micro-nozzle tip and the Lift heads can be connected to the adapter.

The connection is made with a simple push/pull movement.




ADAPTER CONNECTION


AUXILIARY HEAD CONNECTION


MICRO-HEAD CONNECTION


LIFT HEAD CONNECTION

→ POWER CORDS

Your equipment comes with two power cords with different plugs.
Before powering on your device, first connect each of the two power cords to their respective base:

C13 Plug: to the table base

C19 Plug: to the box base

Next, connect each of other ends to a standard grounded electrical wall socket.

TABLE POWER CODE CONNECTION

BOX POWER CODE CONNECTION

If the power cords for your device are damaged, please contact LPG Systems Customer Service for replacements.

LPG Systems Customer Service:
+33 (0)4 75 78 69 89

↗ ATTENTION

The cord with the rectangular (C19) plug must be connected to the box.

→ MAINTENANCE LOG SHEET

Replacement of filter cartridges: To be done when the warning message appears.

Replacement of Endermolift Kit: To be done when the flaps no longer treat the skin properly. They should be replaced after approximately every **14 hours** of use.

DATE	NO. OF HOURS	OPERATIONS PERFORMED

↗ ATTENTION

The device does not require calibration

→ WHAT IFI HAVE A PROBLEM?

If your unit is not working properly, proceed with the following checks before calling Customer Services:

- Is the device properly connected to a power supply plug?
- Is the power supply plug on?
- Is the ON switch lit up?
- Are the filter cartridges clean and correctly placed?
- Is the hose free of obstruction?
- Is the hose properly connected?
- Are the sealing valves of the main head correctly placed, clean and moving?
- Is the Keymodule of the main head correctly fitted?

Once these checks have been carried out and if the malfunction persists, please contact Customer Services of LPG Systems or the nearest authorized dealer, indicating the model of your unit and its serial number.

LPG Systems Customer Service:
+33 (0)4 75 78 69 89

→ TECHNICAL SPECIFICATIONS

Dimensions; LxWxH . . . . . 1850 x 700 x 1490 mm
Net weight . . . . . 140 kg
Maximum set depression. . . . . 69 kPa (690 mbar)
Cooling. . . . . by mechanical ventilation incorporated in the pump
Protection index . . . . . IP 20
Electrical protection class 1
Operating temperature . . . . . +10 °C to +30 °C
Storage temperature . . . . . -20° C to +70° C
Box electrical features . . . . . 100-230V / 50-60Hz / 650-625W

Table electrical features

Voltage (V)	230	120	100	100
Frequency (Hz)	50	50	50	60
Intensity consumed (A)	1	1,8	2,1	2,1

Operating environment:
Ambient temperature:. . . . . 10 °C to 30 °C for normal operation.
Ambient relative humidity: . . . . . 30% to 75% without condensation.
Atmospheric pressure: . . . . . no significant influence for operation.
Max altitude: . . . . . 2,500 m

General characteristics:
Adjustable height: 65 cm to 90 cm from ground
Back adjustment: 0° to 70°/from horizontal

Unit fitted with patented treatment heads.

ELECTROMAGNETIC COMPATIBILITY
For more information about electromagnetic compatibility, refer to the "Electromagnetic Compatibility" appendix.

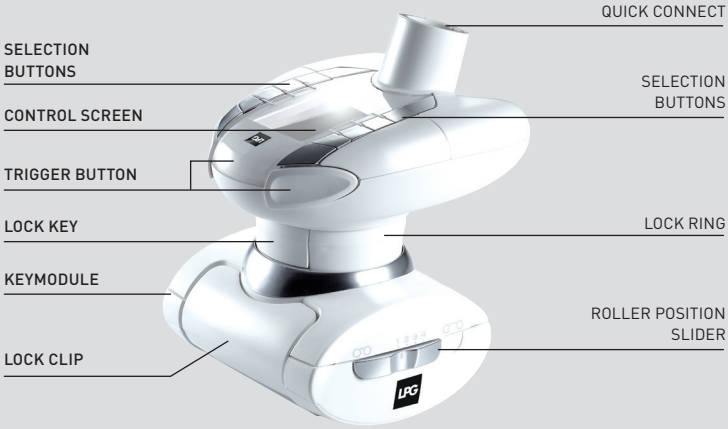
The Cellu M6® Endermolab i device is marked CE 1912 as a medical device by virtue of Annex II of regulation 93/42/EEC (applicable standards IEC 60601-1 Ed3 and related standards)

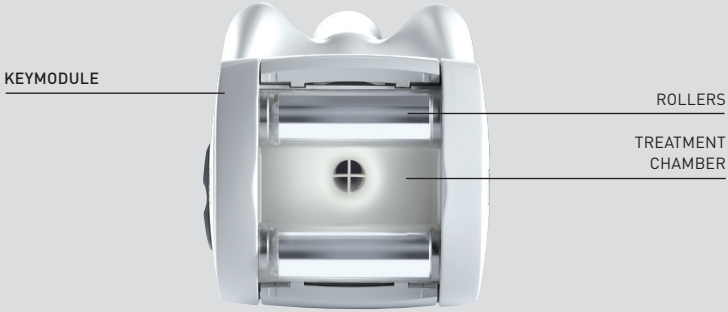
Cellu M6® Endermolab

TREATMENT HEADS

→ INDEX

→ ERGODRIVE™ HEAD DESCRIPTION





→ KEYMODULE™ SET DESCRIPTION

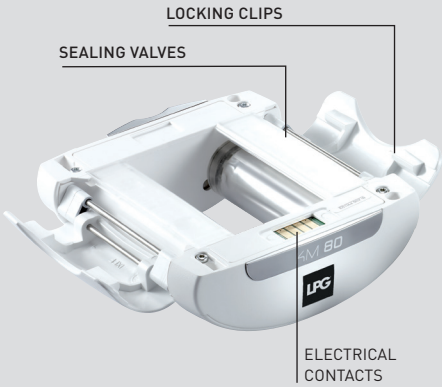
The Keymodule™ set is the interchangeable lower section of the Ergodrive head. It comprises the motorised rollers and sealing valves.

You have one Keymodule™ set (KM80).

Installation:

The Keymodule™ set is designed to be compatible with the treatment head. Attach the Keymodule™ set as shown below.

To remove the Keymodule™ set release the latches.



→ SLIDER SETTINGS

The Keymodule has a slider which can be adjusted to four positions, making it possible to adjust the spacing between the motorised rollers.

When the slider is positioned to the left **(Fig. 1)**, this allows the maximum level of mobility in the rollers.

When the slider is positioned to the right **(Fig. 2),** the rollers cannot move.

When the slider is in an intermediate position, the rollers have reduced mobility.

FIG. 1

FIG. 2

→ FREE ROTATION FUNCTION

The Ergodrive head includes a function that allows the Keymodule to rotate freely.

To do this, lift the lock ring upward until it clicks into position **(Fig. 1- 2).**

The head can now rotate freely **(Fig. 3)**.

To lower the lock ring and lock the postion of the head, press the lock key **(Fig. 4).**


FIG. 1


FIG. 2


FIG. 3


FIG. 4

The head rotation can be locked in any of the four positions below:


ROTATION 0


ROTATION 1


ROTATION 2


ROTATION 3

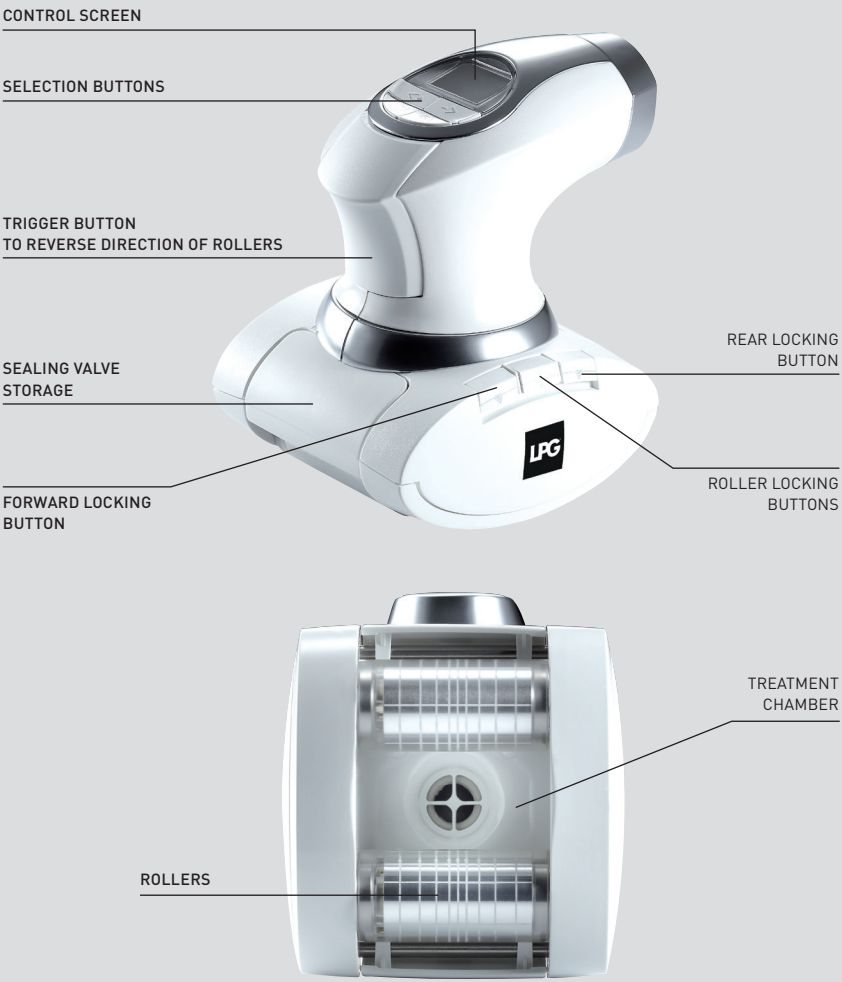

ROTATION 4

These four positions can be found by:

a) following the preceding instructions for free rotation, then pressing the lock key and rotating the head until it clicks into the desired position.

b) lifting the lock ring halfway and then rotating the head to the desired position until it clicks into place.

→ TR50 HEAD DESCRIPTION



→ TR50 HEAD DESCRIPTION (CONT'D)

Locking the rollers:

The rollers on the TR50 can be locked simply by pressing the appropriate buttons, as shown in the photos:

ROLLERS UNLOCKED

FRONT ROLLER LOCKED - FAR

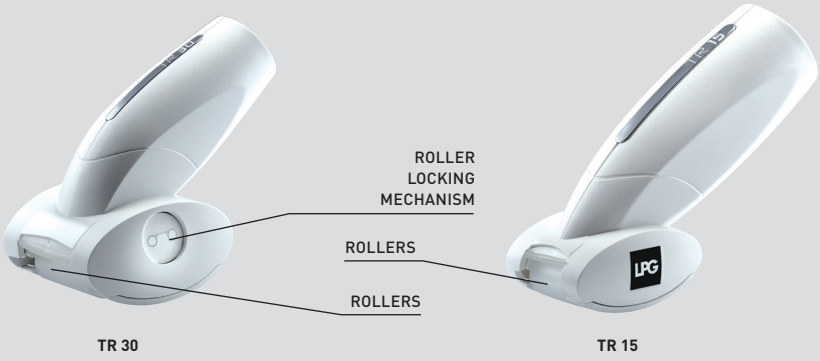
REAR ROLLER LOCKED - FAR

REAR ROLLER LOCKED - NEAR

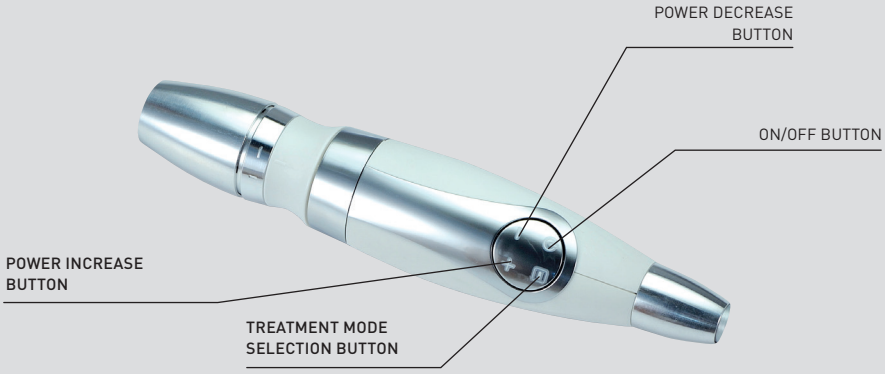
FRONT AND REAR ROLLERS LOCKED - NEAR

Reversing the Roller Direction:
The direction of the rollers reverses each time the trigger is pressed.

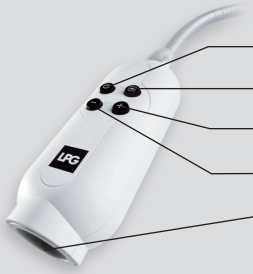
→ AUXILIARY HEADS DESCRIPTION



→ ADAPTER DESCRIPTION



→ ERGOLIFT™ HEAD DESCRIPTION



ON/OFF BUTTON

SELECTION BUTTON FOR TREATMENT TYPE

POWER INCREASE BUTTON

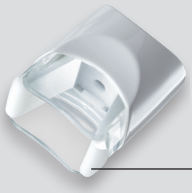
POWER DECREASE BUTTON

HOSE CONNECTION

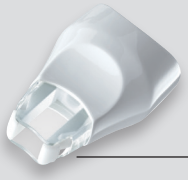
ERGOLIFT CHAMBER CONNECTION

They can be connected and disconnected by a simple push-pull action.

→ ERGOLIFT™ CHAMBER DESCRIPTION

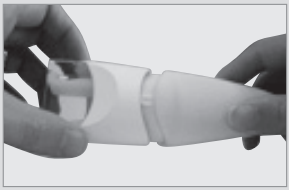


Lift 20
Treatment chamber with removable flap



Lift 10
Treatment chamber with removable flap

Only the LIFT20 and LIFT10 can be connected to the Ergolift™ head. They can be connected and disconnected by a simple push-pull action.



→ LIFT HEADS DESCRIPTION

Adjusting the flap:

To ensure that the flap retains its position, thereby optimizing performance, it is important to adjust the dial settings according to the treatment intensity:

• Turn the dial from A to E to increase the flap's return force.

• Turn the dial from E to A to decrease the flap's return force.

TML10
WITH REMOVABLE FLAPS

TML20
WITH REMOVABLE FLAPS

TML30
WITH REMOVABLE FLAPS

→ MICRO-NOZZLES AND MICRO-HEADS DESCRIPTION

→ CLEANING THE ERGOLIFT™ HEAD AND ERGOLIFT™ CHAMBERS

For hygienic reasons, the treatment heads should be cleaned after each use, using antiseptic wipes soaked with a bactericidal and fungicidal solution.
Special attention must be given to the cleanliness of the parts that are in contact with the patient.

Before each use, clean the flap and Ergolift™ chamber:

1. Disconnect the chamber from the Ergolift™ treatment head. **(Fig. 1)**
2. Remove the flap thanks to the dedicated tool. **(Fig. 2)**
3. Thoroughly clean the Ergolift™ chamber, the flap and the tool for at least one minute with the wipes as describe here below. **(Fig. 3)**
4. Put the flap back in the Ergolift™ chamber by following the same steps in the reverse order **(Fig. 4)**

FIG. 1 REMOVAL TOOLS

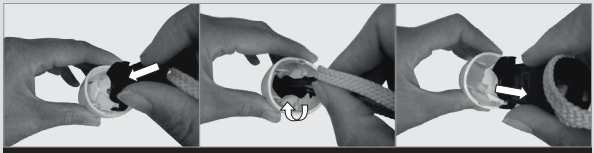


FIG. 2 ROTATION 90°

→ CLEANING THE ERGOLIFT™ HEAD AND ERGOLIFT CHAMBERS (CONT'D)

FIG. 3

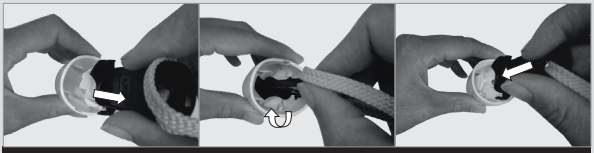


FIG. 4 ROTATION 90°

→ DISINFECTING THE ERGOLIFT™ CHAMBERS

The Ergolift™ head is in direct contact with patient's skin. Under certain specific applications, it needs to be disinfected after each use.

1. Follow the cleaning procedure described above.

2. Soak the flap and Ergolift™ chamber in an OPA disinfectant for 12 minutes at 20 °C, as recommended on the disinfectant packaging.

3. Carefully rinse the flap and the Ergolifttm chamber with sterile or drinking water for at least one minute using a large volume of water (approximately eight litres). Repeat twice for a total of three rinses..

4. Dry the Ergolift™ chamber and flap.

5. Clean the storage drawer with antiseptic wipes, then place the Ergolift™ chamber and flap in it.

↗ ATTENTION

The use of aggressive products, such as acetone, trichloroethylene or alcohol at 90° and abrasive sponges, ultrasound or UV lamps is strictly prohibited. All cleaned and/or disinfected heads should be placed in the storage drawer to avoid any confusion. Use a disinfectant whose active ingredient is ortho-phthalaldehyde (OPA). Before using the disinfectant, read and follow the recommendations, contraindications and warnings associated with this product. Refer to the instructions for using this solution. All the procedures described in this section must be carried out with the machine turned off and the power cord unplugged.

→ REMOVING THE TML20 AND TML30 FLAPS

1. Take the Lift head and insert the beveled side of the correct removal tool into the opening between the flaps.

2. Pull out the tool and the flaps will come out with it.

3. Remove the flaps from the tool and clean them carefully with a wipe.

4. Take the smooth side of the tool and place the flaps back into position on the tool.

5. Insert the flaps back into the treatment head until they are in the correct position.

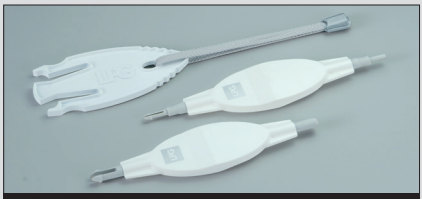

REMOVAL TOOLS


FIG. 1


FIG. 2


FIG. 3


FIG. 4


FIG. 5

→ REMOVING AND CLEANING THE TML10 FLAPS

1. Take the Lift head and insert the beveled side of the correct removal tool into the opening between the flaps.

2. Pull out the tool and the flaps will come out with it.

3. Remove the flaps from the tool and clean them carefully with a wipe.

4. Take the smooth side of the tool and place the flaps back into position on the tool.

5. Insert the flaps back into the treatment head until they are in the correct position.

FIG. 1

FIG. 2

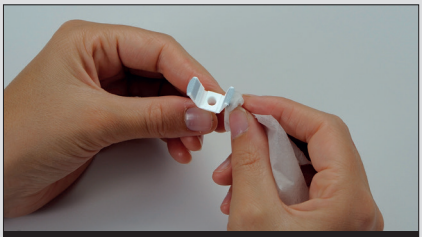


FIG. 3

FIG. 4

FIG. 5

→ CLEANING THE ERGODRIVE™ HEAD

Cleaning the sealing valves:

1. Release the Keymodule™ **(Fig. 1)**.

2. Move the rollers to the center.

3. Remove the corresponding sealing valve by handling it as shown on the photo. Repeat the operation for the other valve **(Fig. 2)**.

4. Meticulously clean the flaps and their housing for at least one minute using LPG wipes soaked in a bactericide and fungicide solution **(Fig. 3-4).**

Replace the sealing valves after **100 hours** operation.

FIG. 1

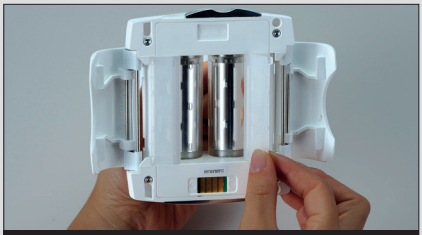


FIG. 2


FIG. 3


FIG. 4

→ CLEANING THE ERGODRIVE™ HEAD (CONT'D)

Cleaning the Keymodule™ set:

1. Turn over the Keymodule™ set and thoroughly clean the following parts for at least one minute with the LPG® wipes soaked in a bactericide and fungicide solution:
   a) The housing on both sides of the rollers **(Fig. 1).**
   b) The rollers (rotate them manually to clean the entire surface) **(Fig. 2).**
   c) The sealing ring between the Keymodule™ set and head housing.
   d) The plastic covering on the Keymodule™ set.

2. Turn the Keymodule again and refit the sealing valves.

3. Ensure that the sealing joint between the Keymodule™ set and main head is also cleaned.

4. Check that the electrical contacts are clean and dry and proceed in the reverse sequence to refit the Keymodule™ set onto the main head.

5. Clean the storage drawer using LPG® wipes, then place the head in it.


FIG. 1


FIG. 2


FIG. 3


FIG. 4

→ CLEANING THE TR50 HEAD

1. Remove the sealing flaps as shown in the photos below **(Fig. 1 to 4)**.
2. Thoroughly clean for at least one minute using LPG wipes soaked in a bactericide and fungicide solution:
a) Flaps and their housing **(Fig. 5 and 6)**
b) The casing on both sides of the rollers (turn the head over, rotate the rollers manually to clean the entire surface) **(Fig. 7 to 10)**
c) The sabot
3. Reattach the sealing flaps.
4. Clean the storage drawer using LPG® wipes, then place the head in it.

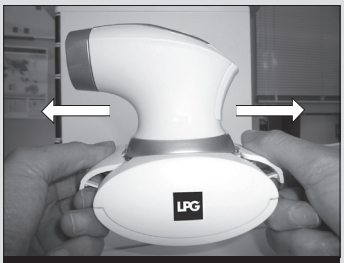


FIG. 1

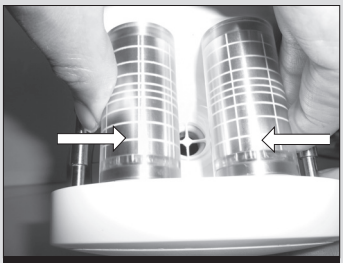


FIG. 2

FIG. 3

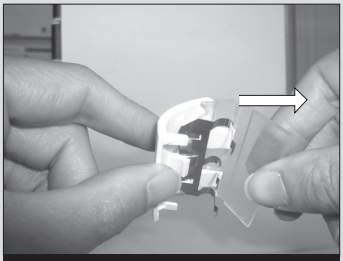


FIG. 4

→ CLEANING THE TR50 HEAD (CONT'D)


FIG. 5


FIG. 6

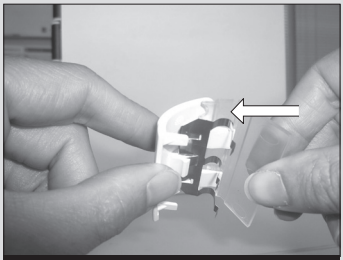


FIG. 7

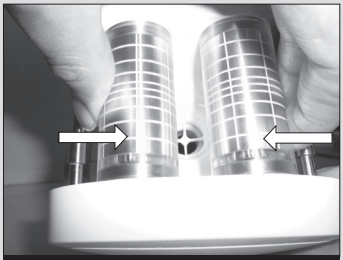


FIG. 8

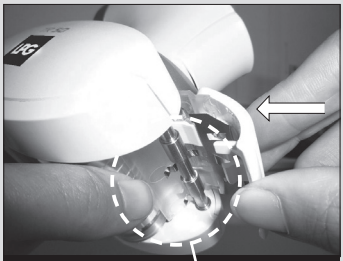
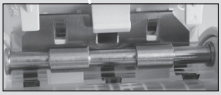
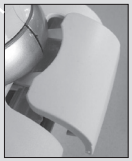


FIG. 9


FIG. 10





→ CLEANING THE AUXILIARY HEADS, MICRO-HEADS AND MICRO-NOZZLES

1. Disconnect the auxiliary head from the adapter.

2. Remove the two rollers from the head for effective and rapid cleaning **(Fig. 1-2).**

3. For the auxiliary heads use the dedicated tool provided **(Fig. 3 to 4)**.

4. Thoroughly clean for at least one minute the rollers, seal, treatment chamber, micro-heads, disassembly tool and micro-nozzles with LPG wipes soaked in a bactericide and fungicide solution **(Fig. 5 and 6).**

5. Refit the rollers and check they spin freely.

6. To clean the auxiliary heads, use cotton soaked with the same solution.

7. Clean the storage drawer using LPG wipes, then place the heads in it.

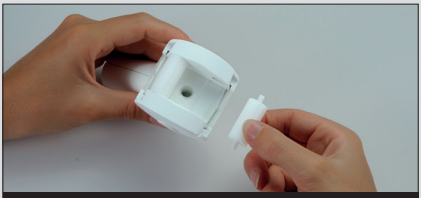


FIG. 1


FIG. 2


FIG. 3


FIG. 4


FIG. 5


FIG. 6

→ DISINFECTING THE AUXILIARY HEADS

The motorised treatment heads (Ergodrive™ and TR50) need to be used with an Endermowear™ suit. Non-motorised treatment heads (auxiliary heads, micro-nozzles and micro-heads) can be used directly on the skin in specific cases.
In these cases, the heads need to be disinfected after each use:

1. Use the cleaning procedure described above.

2. Soak the rollers, micro-heads, disassembly tool and micro-nozzles in a disinfectant for 12 minutes at 20 °C, as recommend on the disinfectant packaging.

3. Carefully rinse the flap and the treatment chamber with sterile or drinking water for at least one minute using a large volume of water (approximately eight litres). Repeat twice for a total of three rinses.

4. Dry the parts.

5. Pre-clean the storage drawer using LPG® wipes, then place the head in it.

→ REPLACING THE KEYMODULE™ SET

The Keymodule™ set is a high-technology unit comprising numerous mobile micro-mechanical parts.
The unit must be returned to our Technical Support Center if certain worn items require replacing.

The head may start to lose some of its characteristics at approximately **1000 hours** of operation.

↗ ATTENTION

Use a disinfectant whose active ingredient is ortho-phthalaldehyde (OPA). Before using the disinfectant, read and follow the recommendations, contraindications and warnings associated with this product. Refer to the instructions for using this solution. All the procedures described in this section must be carried out with the machine turned off and the power cord unplugged. - Do not use corrosive products such as acetone, trichloroethylene or rubbing alcohol, nor abrasive sponges.

→ ENDERMOWEAR

LPG Endermowear™ suit, available in several sizes for both men and women, is a clothing specially designed for superior treatment of the entire body. For personal use, it ensures perfect hygiene and its opaque areas protect client's privacy during the treatment. The Endermowear unique composition ensures an excellent grip of the skin for facilitating smoother movements of the treatment head as well as protecting it.

The products come with a re-enforced bag with empty client name tag for privacy. Once personalized, it becomes the customer's property and can be a multisession-use garment requiring the client to wash it between each session. Require the client to check and follow the washing instructions printed on bag's tag.

→ GENERAL WARRANTY CONDITIONS

You have recently acquired a device distributed by LPG Systems or an LPG Systems approved distributor. It is the purchaser/ user's responsibility to find out from the local authorities the conditions and professional qualifications required before using the appliance.

The purchase of this equipment implies the legal acceptance by the purchaser/ professional user of these general warranty conditions. If the appliance was sold to you by an approved LPG Systems distributor, the purchaser/ user should refer to the supplier's warranty conditions. These may in no way increase the undertakings made by LPG Systems in these present warranty conditions.
The warranty can only be implemented and is only valid if the warranty slip has been duly filled out and returned to LPG Systems within two weeks of delivery, irrespective of the country. Warranty slips that are only partially completed will be rejected. The appliance is guaranteed against manufacturing flaws and defects in the raw materials.

The warranty extends for the shorter of the following two periods: two (2) years OR two thousand (2000) hours of use from the invoice date. During this period, LPG Systems undertakes to exchange or repair free of charge, as quickly as possible, any part that LPG Systems acknowledges as defective; however LPG Systems does not undertake to replace the entire appliance.

Traveling and living expenses for our technicians and transportation costs of the appliance or parts to and from the aftersales service workshop are not covered by this warranty. Replacements and repairs performed within this warranty, with or without immobilization of the equipment, shall not have the effect of extending the warranty period.

Replaced parts become the property of LPG Systems or the approved distributor. No compensation shall be paid for loss of use. Subject to other conditions hereafter, warranty shall apply if the purchaser/professional user has allowed LPG Systems to proceed to necessary repair works.

→ GENERAL WARRANTY CONDITIONS (CONT'D)

Warranty exclusion:

- Damage occurring during transportation. Transportation of this equipment and/or spare parts is at the recipient's own risk. Before accepting delivery, it is the recipient's responsibility to verify the state of the goods and to make a claim against the transport company in the manner usual in the delivery country.
- Non-observance of the installation and operating instructions, failure to carry out maintenance and/or negligence in maintaining the appliance and/or its filter cartridges, connection to a faulty electricity supply or a non-grounded electricity supply or a power supply whose voltage is different to the one indicated on the appliance.
- Modification, mounting of accessories or dismantling of the equipment.
- Any operation and/or intervention not specified in the LPG Systems Operating Instructions and performed on the equipment by the purchaser/user and/or any party not approved by LPG Systems.
- Use of consumables, spare parts, inappropriate components or parts not supplied by LPG Systems.
- Blockage of the appliance through aspiration of a foreign body.
- Normal wear and tear of any of the equipment's parts resulting from normal usage.
- Damages or default resulting of accidental events (falls, impacts, etc.)
  Damages or default resulting of natural disasters (lightening, water damages, etc.)
  Fire, negligence or abuses.

If an appliance is sold before the end of the warranty period, the warranty is transferred to the purchaser for the remaining warranty period, on the condition that:
I- The original invoice is provided.
II- That the initial vendor is informed of the sale.

→ LIMITATION AND EXONERATION OF LIABILITY

The warranty is limited to the replacement of the components of the device which comply with conditions described above. Under no circumstances shall LPG Systems be liable for any loss or damage as a result or in connection with the device and/or its use, including any financial loss, loss of margin, loss of use, etc. This clause shall apply under any and all legal basis. Whenever the above restriction may not be applicable or enforceable, LPG Systems' liability shall be limited to the price for the device and/or the service.

Failure to comply with the general warranty conditions during the warranty period and after its expiry may constitute an exonerating cause of liability of LPG Systems in case of damage attributable to the delivered products.

The purchaser/user is responsible for the use of the device and will assume full responsibility for any damages, including damages caused to third parties, resulting from the failure to observe the instructions for use of the device and/or resulting from an improper use.

Under no circumstances LPG Systems will be held liable for any intangible or indirect damages, including any commercial or financial loss, loss of profit, loss of earnings and damage to the brand image.

LPG Systems' liability, for all causes (with the exception of personal injury), is limited to the amount of the defective device's price.

The purchaser/user is solely liable for his prescriptions, care and information to his customers/patients. The responsibility of the delivery of care by the purchaser/user within his structure is held by him and is subject to his sole discretion.

By consequence, LPG Systems will in no case be held liable in event of inappropriate:
1- Use of the device
2- Prescription
3- Protocol
4- Care and any contraindications not respected

→ WARRANTY ACTIVATION

You can activate your warranty online by connecting to our warranty webpage:

http://warranty.lpgsystems.com

→ APPENDIX: ELECTROMAGNETIC COMPATIBILITY

TABLE 1: DIRECTIVES AND MANUFACTURER DECLARATION – ELECTROMAGNETIC EMISSIONS		
The CELLU M6® ENDERMOLAB *i* device is intended for use in the electromagnetic environment specified below. The CELLU M6® ENDERMOLAB *i* customer or patient should ensure that it is used in such an environment.		
Emissions test	**Conformity**	**Electromagnetic Environment - Directives**
RF emissions CISPR 11	Group 1	The CELLU M6® ENDERMOLAB *i* device uses RF energy only for its internal functions. Therefore, its RF emissions are very low and unlikely to cause interference in nearby electronic devices.
RF emissions CISPR 11	Class B	The CELLU M6® ENDERMOLAB *i* device may be used in all establishments, including domestic sites and sites that are directly connected to the low voltage public power grid, which supplies domestic buildings.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations and flicker IEC 61000-3-3	Conforms	

TABLE 2: IMMUNITY				
Test	**Requirements**		**Level of conformity**	
Electrostatic discharge (DES) IEC 61000-4-2	± 8 kV on contact ± 2/4/8/15 kV in the air		± 8 kV on contact ± 2/4/8/15 kV in the air	
Radiated RF electromagnetic fields IEC 61000-4-3	10V/m 80MHz-2.7 GHz 80% AM to 1 kHz		10V/m 80MHz-2.7 GHz 80% AM to 1 kHz	
Proximity fields issued by RF wireless communication devices IEC 61000-4-3	**Frequency (MHz)**	**Modulation**	**Requirements (V/m)**	**Conformity (V/m)**
	385	Pulsed modulation: 18 Hz	27	27
	450	Pulsed modulation: 18 Hz	28	28
	710 – 745 - 780	Pulsed modulation: 217 Hz	9	9
	810 – 870 - 930	Pulsed modulation: 18 Hz	28	28
	1720 – 1845 - 1970	Pulsed modulation 217 Hz	28	28
	2450	Pulsed modulation 217 Hz	28	28
	5240 – 5500 - 5785	Pulsed modulation 217 Hz	9	9
Fats transient / burst IEC 61000-4-4	Power lines: ± 2 kV Input/output lines: ± 1 kV Repetition frequency: 100 kHz		Power lines: ± 2 kV Input/output lines: ± 1 kV Repetition frequency: 100 kHz	
Surges IEC 61000-4-5	Between phases: ± 0.5 kV, ± 1 kV Between phases and ground ± 0.5 kV, ± 1 kV, ± 2 kV		Between phases ± 0.5 kV, ± 1 kV Between phases and ground ± 0.5 kV, ± 1 kV, ± 2 kV	
Conducted RF disturbances IEC 61000-4-6	3 V 0.15 MHz – 80 MHz 6 V in ISM bands and amateur radio bands between 0.15 MHz and 80 MHz 80% AM to 1 kHz		3 V 0.15 MHz – 80 MHz 6 V in ISM bands and amateur radio bands between 0.15 MHz and 80 MHz 80% AM to 1 kHz	
Power frequency magnetic field IEC 61000-4-8	30 A/m		30 A/m	
Voltage Dips and Interruptions: IEC 61000-4-11	0% UT; 0.5 cycle A 0°, 45°, 90°, 135°, 180°, 225°, 270° et 315° 0% UT; 1 cycle to 0° 70% UT; 25/30 cycles to 0° 0% UT; 250/300 cycles		0% UT; 0.5 cycle A 0°, 45°, 90°, 135°, 180°, 225°, 270° et 315° 0% UT; 1 cycle to 0° 70% UT; 25/30 cycles to 0° 0% UT; 250/300 cycles	

→ APPENDIX: ELECTROMAGNETIC COMPATIBILITY

Conducted RF disturbances IEC 61000-4-6	3 V 0.15 MHz – 80 MHz 6 V in ISM bands and amateur radio bands between 0.15 MHz and 80 MHz 80% AM to 1 kHz	3 V 0.15 MHz – 80 MHz 6 V in ISM bands and amateur radio bands between 0.15 MHz and 80 MHz 80% AM to 1 kHz
Power frequency magnetic field IEC 61000-4-8	30A/m	30 A/m
Voltage Dips and Interruptions: IEC 61000-4-11	0% UT; 0.5 cycle A 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% UT; 1 cycle to 0° 70% UT; 25/30 cycles to 0° 0% UT; 250/300 cycles	0% UT; 0.5 cycle A 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0% UT; 1 cycle to 0° 70% UT; 25/30 cycles to 0° 0% UT; 250/300 cycles

Your Cellu M6® Endermolab i device requires special care concerning the EMC; it must be installed and service according to the information provided in this user guide.

Portable and mobile RF communication devices must not be used within 30 cm of your device, they can cause undesired operation.

The use of other treatment heads than those provided by LPG Systems may result in increased emissions or decreased immunity of the device.

Your Cellu M6® Endermolab i device should not be used adjacent to or stacked with other medical devices.

The Cellu M6® Endermolab i device does not manage essential performances.

Interference may occur near equipment marked with the following symbol: 

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