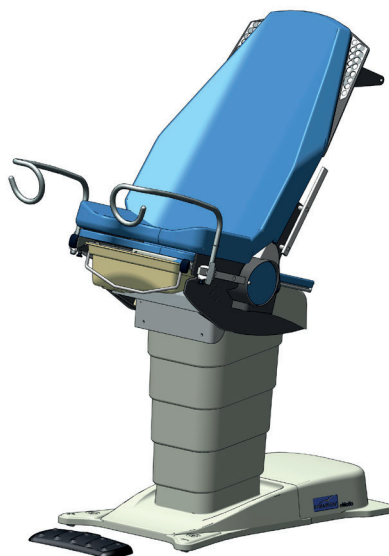


User's Guide



Model Shown : 15640-01



Model Shown : 15620-01



Electric Gynecological Table
15620-01 / 15624-01 / 15626-01
15630-01 / 15640-01

Product Identification

Model Number :

Serial Number :

Date of Purchase :

Name of Owner / Facility :

Name of Dealer :

Dealer's Phone Number :

Promotal Authorized Service Company :

Legal Notice

PROMOTAL

22, rue de Saint-Denis de Gastines
B.P. 26 - 53500 ERNÉE Cedex
FRANCE

Tél. : +33 (0)2 43 05 12 70

Fax : +33 (0)2 43 05 68 99

internet : www.promotal.com

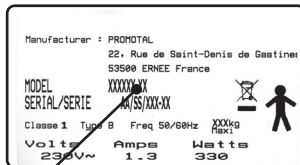
The descriptions and specifications contained in this

Operating Manual are deemed correct at the time of printing.

Promotal, however, reserves the right to modify its models and its procedures or render them obsolete without notice.



Before any order, we recommend that our customers consult a local sales manager.

TYPE PLATE



Year (YY)/Week (WW) of manufacture



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Important information

Safety symbols



Warning sign

Information marked with this symbol must be read and strictly complied with!

Remark

Draws your attention to a procedure, practice or situation.



Humidity limitation



Maximum number of stacked pallets



Correct orientation for transportation



Atmospheric pressure limitation



Temperature limitation



Fragile



Keep in a dry place



Not to use sharp objects



Manufacturer



Type B applied part



Fuse rating



Protective earth



Dangerous voltage / risk of electrocution



To be disposed off separately from household waste



Equipotential terminal



Life cycle (motor cycle)
1 minute working / 9 minutes stopped



General safety sign



Housing protection classification



Catalog reference



serial number



For indoor use only



Refer to the instruction manual / brochure



CE marking

Remark

the equipotential terminal allows the connection of an equipotential conductor to bring all the different elements of an electro-medical system to the same potential.



Do not sit on the back rest



ban on using the footrest as a step



You must not use the medical device for transfers

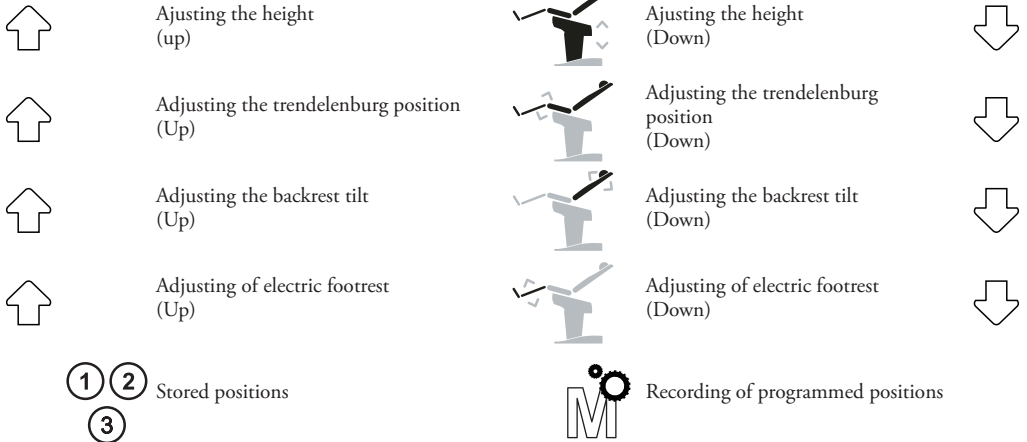


Prohibition to sit on the leg rest



Warning

You must not remove the pictograms and warning signs provided by the manufacturer! The manufacturer disclaims all responsibility in case of removal of these signs.



Applied parts

The applied parts according to standard EN 60601-1 are:

- PVC upholstery
- Leg rests / Leg supports
- Gynecological examination stirrups (accessories)

Electrical power supply



The equipotential terminal must not be used as a protective earth connection under any circumstances.



This medical device has electrical classification 1, it must only be connected to a power supply equipped with a protective earth.

Electromagnetic interference

This Promotal medical device was designed and built to minimize electromagnetic interference with other equipment. If interference is, however, observed, you must remove the apparatus causing the interference from the room and/or plug it into an isolated circuit.

Unpacking precautions

Medical device delivered on a wooden pallet

The medical device positioned on a wooden pallet may be easily moved using a forklift truck, as long as this is used correctly. Before transportation,

ensure that the forklift truck is correctly positioned in relation to the pallet, and that the unit is stable.

Storage conditions

Room temperature:

Relative humidity :

Atmospheric pressure :

-15 °C to +60 °C (+5 °F to 140 °F)

10 % to 90 % (without condensation)

0.5 bar to 1.05 bar (500 hPa to 1050 hPa)

All storage must be carried out in accordance with the following recommendations:

- Clean, aired and temperate area.
- Medical device stored in an area sheltered from bad weather and direct sunlight.
- Dry room.
- Medical device protected from shocks.

- Do not store in an area subject to frequent passage.
- Do not stack material.
- Keep in its original packaging until the final destination.

Conditions of use

- Dry and temperate area.
- Maximum altitude : 2000 m

- Temperature 10 to 40° C
- Relative humidity 75% maxi.

Unpacking and Installation


Step by step

1) During unpacking, remove all staples and remove the cardboard packaging carefully.

Caution : be careful with cutting tools, as fragile parts of the medical device (*covering, plastic housing, etc.*) may be near them.

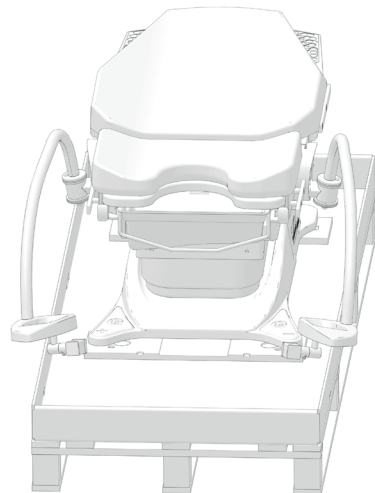
2) If possible, transport the medical device on its pallet up to the final place of use.

The medical device is adjusted to a resistant position for transportation.

see: as indicated in the diagram 

3) Cut the two green bands that fix the DM to the pallet..

4) Next, take the DM off the pallet.



Caution

Four people are required to remove the medical device from the pallet.

Check

Having unpacked the medical device, follow these steps:

- 1) Check the delivery documents to ensure that the delivery is complete.
- 2) Check the external components for any damage during transportation.

3) Check that the packaging contains the medical device, accessories and Options, the supply cable (*if electrical MD*) and the User Guide.

Remark:

Within the European Union, all problems, complaints or questions should be addressed to:

Promotal
22, rue de Saint-Denis de Gastines
53500 Ernée,
FRANCE
Telephone : + 33 (0)2 430 517 76
Fax : + 33 (0)2 430 572 00

Cleaning protocol

Warning

It is vital to read the user's manual thoroughly before manipulating this Medical Device. The equipment should only be used for its intended purpose as described in our documentation. Installation and connection must only be carried out by qualified personnel. The electrical components (cylinder, box,

control handle, battery, adapter, etc.) must not be opened under any circumstances. PROMOTAL shall not be held liable for any damage resulting from non-compliance with these instructions.



Any modification to the medical device without written authorization from the manufacturer is forbidden.



Caution

Only accessories designed and provided by Promotal for this medical device are authorised for use.



Caution

This medical device is not intended to be cleaned in a washing tunnel.

Cleaning/Disinfecting

This medical device must be regularly cleaned using the appropriate detergent products and disinfected using bactericidal, virucidal and fungicidal disinfectants.

A mild detergent such as soapy water can be used for routine cleaning of upholstery, stainless steel, aluminium or painted surfaces, plastic parts and control components, followed by effective rinsing and thorough drying.

Detergents and disinfectants designed for use with medical appliances, such as those containing quaternary ammonium compounds, hydrogen peroxide, ethanol, chlorine compounds, etc. can be used on our medical devices provided that:

- The concentration prescribed by the suppliers of such products are complied with;
- The application conditions (contact time, quantity used, temperature, rinsing, etc.) are complied with;
- The supplier's instructions state that the detergent-disinfectant used is suitable for use with:
 - PVC, ABS, Polyamide, Polyurethane, Polypropylene
 - Epoxy-coated metal surfaces
 - Stainless steel or aluminium metallic surfaces.

Warning:

- Solvents are strictly prohibited.
- The use of abrasive powders or any other abrasive product should be avoided.
- High-pressure cleaning is forbidden.



Under no circumstances shall Promotal be held liable under warranty for any damage caused by non-compliance with the use instructions for a detergent-disinfectant.



Intended purpose

This medical device is designed for use in professional premises only:

- Medical office
- Health establishment

This medical device must not be installed in domestic premises.

This medical device is intended for use by health professionals for examinations or gynaecological procedures, in the safest possible conditions.

Patients and companions must not intervene directly on the device. They must not have access to the device's controls.

During the consultation, supervision of children is the responsibility of the parents or of the practitioner.



This medical device must under no circumstances be used as a delivery table.

This medical device must not be used in the presence of anaesthetics flammable in contact with air or nitrous oxide to avoid all risks of explosions.

Patient weight capacity

	= 205 Kg = 450 lbs		= 220 Kg = 480 lbs
---	-----------------------	---	-----------------------

Protection against penetration of liquids

- IP X1

Characteristics

Electric height adjustment from 570 to 1085 mm.

Seat can be lifted by an electric cylinder.

Backrest adjustable by electric cylinder Steel structure with ABS covering.

Electric control by foot pedal and control panel.

Seamless anatomical upholstery, M1 cover.

Integrated paper roll holder (max. width 500 mm - max. Ø 125 mm).

Integrated retractable castor system Adjustable glides under base.

Plastic tray on rail (capacity 6 litres).

Anchoring for colposcope support option.



User manual – gMotio

Electrical connection

This medical device must be connected to the mains supply.

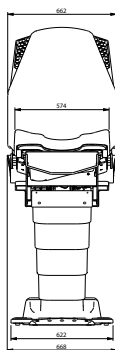
- Connection to the mains supply : 100 V or 240 V (depending on the country)
- Frequency : 50/60 Hz
- Protection classification : Class 1/ type B device
- Absorbed power : 575 W
- Intermittent operating mode : 1 min / 9 min



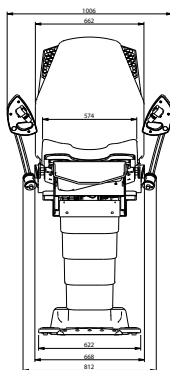
Caution

a power cut could prevent the patient support from being lowered to the low position.
The patient exit must be carried out in the best possible conditions of safety.

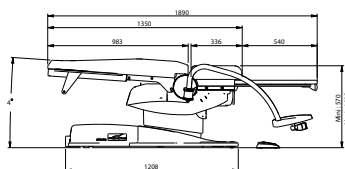
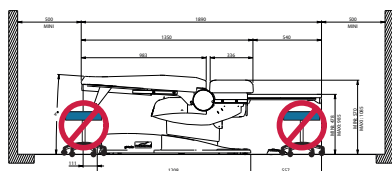
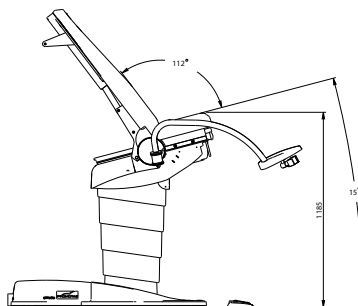
Dimensions / Installation precautions



15620-01 / 15624-01 / 15626-01



15630-01 / 15640-01

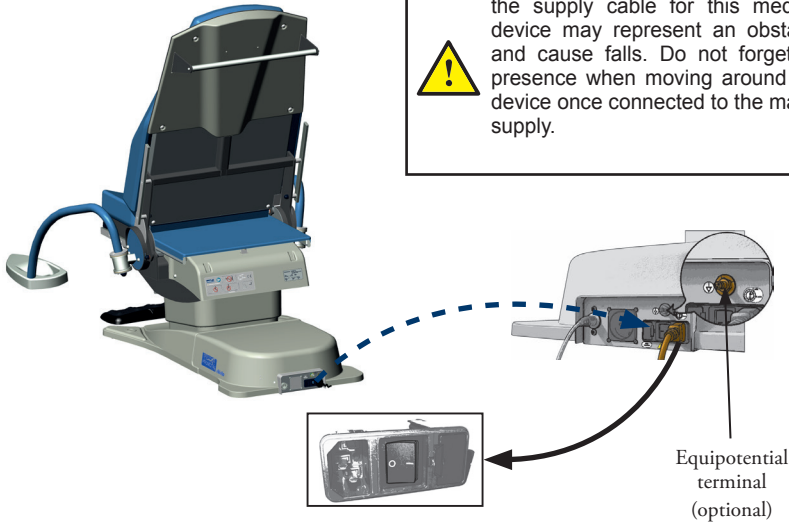
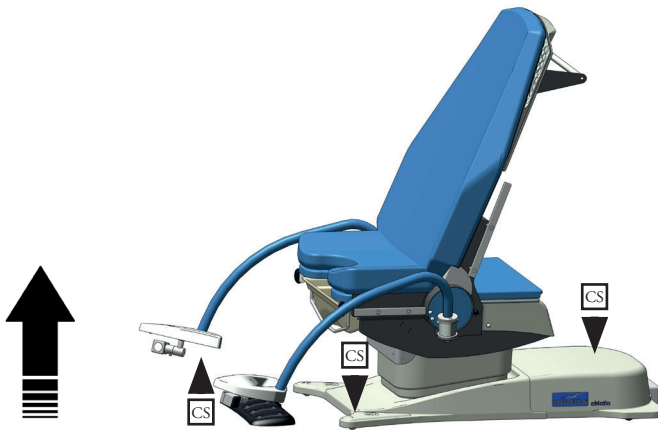


**Caution**

To avoid malfunctions and for safety reasons, no objects must be left under the DM's seat or between its moving frames.

**Caution**

the supply cable for this medical device may represent an obstacle and cause falls. Do not forget its presence when moving around the device once connected to the mains supply.

**Perimeter protection for the user**

In order to avoid any risk of crushing when lowering, the medical device is equipped with sensors (CS). If an object is detected while the medical device is lowering, it will stop lowering, and then rise 5 cm to allow the interfering object to be removed.

**Safety note**

Before using the medical device lowering control, ensure that no objects or obstacles are between the moving parts and the ground.

**Safety note**

When using the position memory function, movements can be interrupted by pushing on one of the command buttons for the DM or, as a last resort, by disconnecting the power supply cord.

**Safety note**

When using the pre-programmed positions, default QE1, QE2, QE3, DM position, DM position), do not leave the patient on the DM without supervision.

**Safety note**

Unplug the power supply cable before moving the MD.

Accumulator

(Built in to the device's power supply system)

(Note: the DM can functions the same in «Battery mode» or in «Mains mode».

Charging the accumulator

Before using the DM for the 1st time, it is recommended to leave it plugged into the mains for 24 hours to enable the accumulator to charge fully.

Note: The DM will lose power after (approximately) 5 to 10 operating cycles in "accu.mode".



Accumulator safety symbols



Certification standard



Do not throw the product away with household waste



Protection rating



Recyclable



Use in dry conditions only



Contains pollutants Lead, etc.



Fuses

Technical characteristics of the accumulator

Voltage	24 V DC
Amperage	1,2 Ah
Accumulator type	Lead gel
Voltage (maximum)	29~45V DC
Charging time	Approx. 8 h (depending on the power supply unit)
Useful life	approx. 1000 cycles (depends on how the DM is used)
Discharge time	approx. 1 an (storage)

Using the individual control

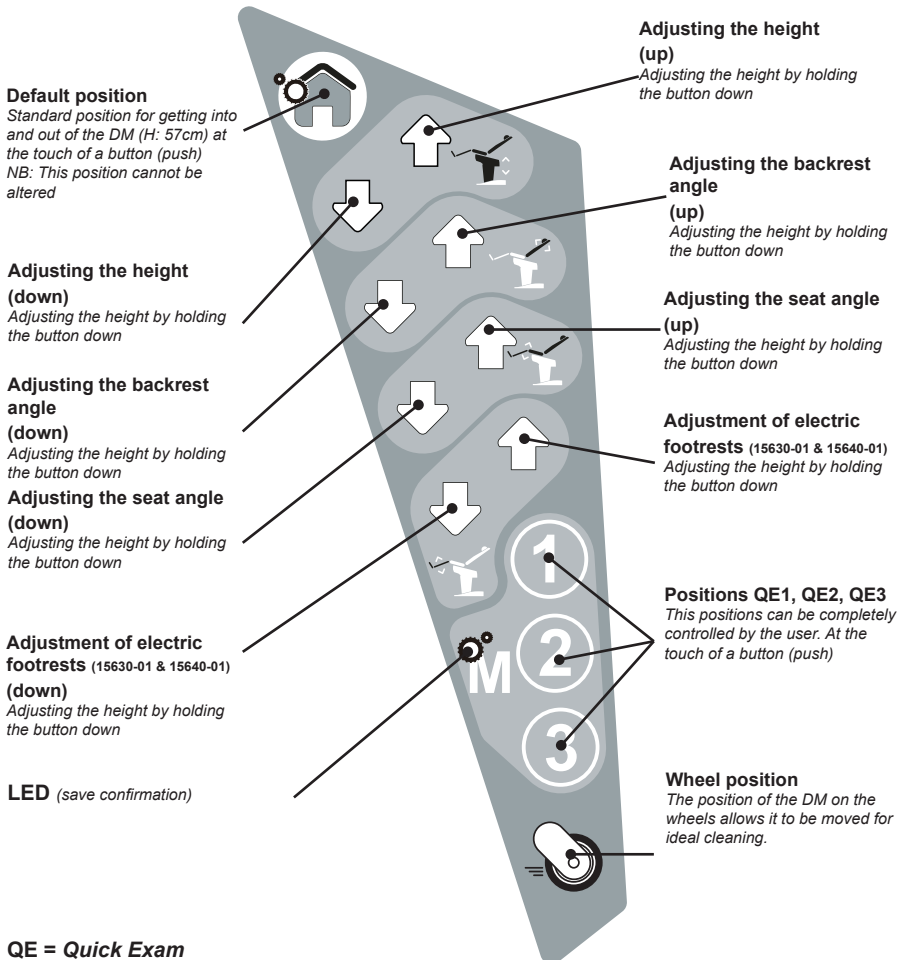
Manual command

Presentation



Safety note

When using the pre-programmed positions, (welcome, QE1, QE2, QE3, DM position, DM position), do not leave the patient on the DM without supervision.



QE = Quick Exam



Safety note

When using the position memory function, movements can be interrupted by pushing on one of the command buttons for the DM or, as a last resort, by disconnecting the power supply cord.

Foot pedal

Technical information :

Type of EI: Radio Frequency (Frequency 2.4G) IEEE802.15.4 standard
Mob.: between 5 and 6 metres

Presentation



Safety note

When using pre-programmed positions (*reception, QE1, QE2, QE3*), do not leave the patient on the DM unattended.

Adjustment of electric footrests (15630-01 & 15640-01)

(up)

Adjusting the height by holding the button down



Adjusting the backrest angle

(up)

Adjusting the height by holding the button down

Adjusting the height

(up)

Adjusting the height by holding the button down

Positions QE1, QE2, QE3

This positions can be completely controlled by the user. At the touch of a button (push)

Adjustment of electric footrests (15630-01 & 15640-01)

(down)

Adjusting the height by holding the button down



Adjusting the backrest angle

(down)

Adjusting the height by holding the button down

Adjusting the height

(down)

Adjusting the height by holding the button down



Default position

Standard position for getting into and out of the DM (H: 57cm) at the touch of a button (push)
NB: This position cannot be altered

QE = Quick Exam



Safety note

When using the position memory function, movements can be interrupted by pushing on one of the command buttons for the DM or, as a last resort, by disconnecting the power supply cord.

Remark: This foot control operates with two (2) LR03 - AAA Alkaline 1.5V batteries.

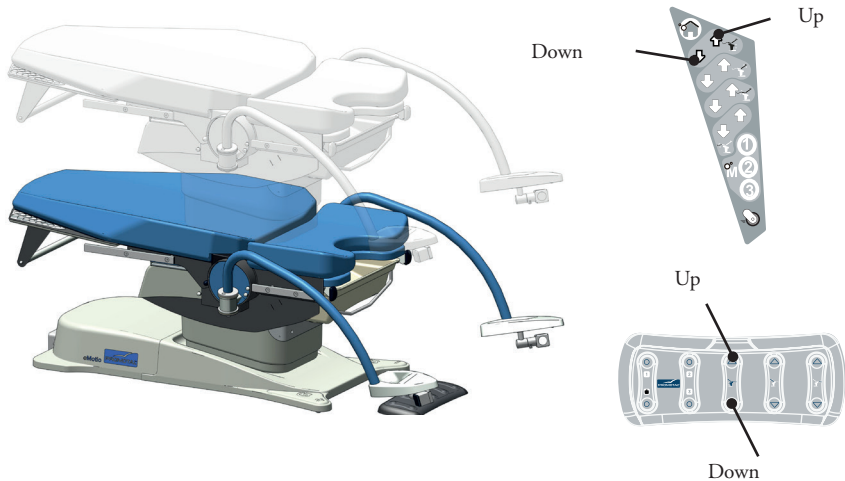
In the event of malfunction, these batteries will have to be replaced.

(See page 48: **REPLACING USED BATTERIES IN THE FOOT PEDAL**)

Using the individual control

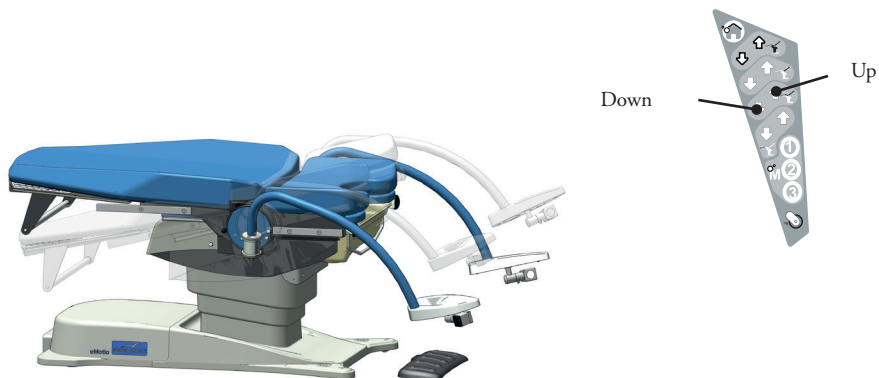
Adjusting the height

The position below is obtained by holding down the hand control or foot control



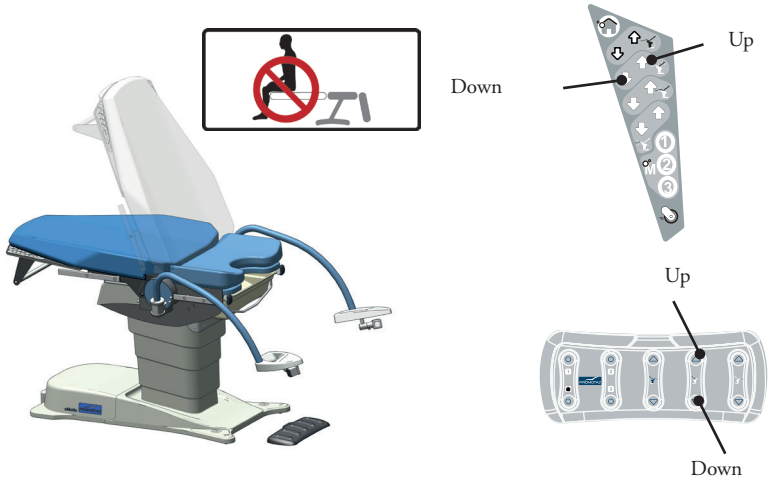
Adjusting the trendelenburg position

The position below is obtained by holding down the hand control or foot control



Adjusting the backrest tilt

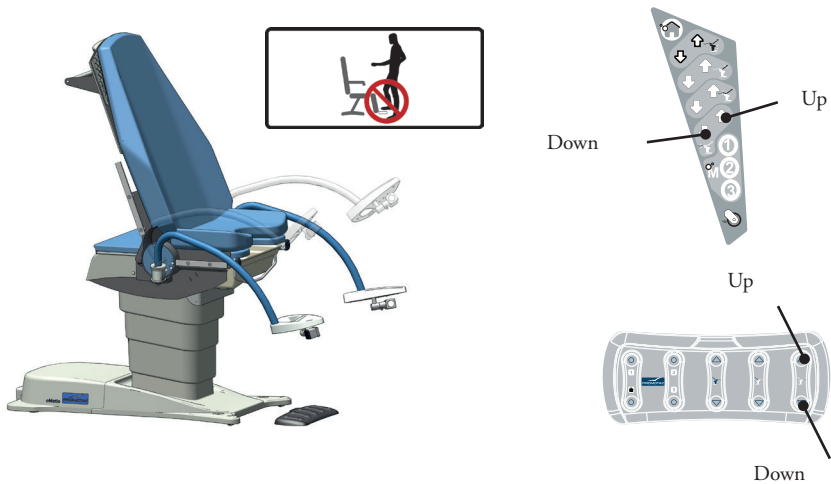
The position below is obtained by holding down the hand control or foot control



Adjustment of electric footrests

(only on models 15630-01 and 15640-01)

The position below is obtained by holding down the hand control or foot control



Safety note



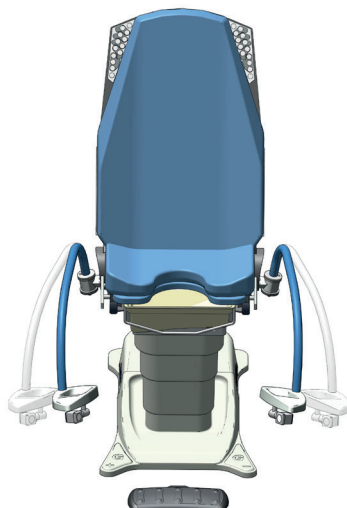
For safety reasons, in order to prevent collisions, the height adjustment of the electric footrests is blocked if the leg rest extension 15672-01 is not fully retracted.

Using the individual control

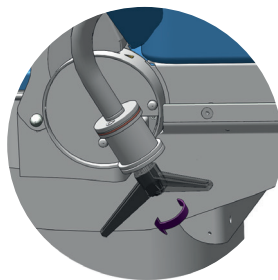
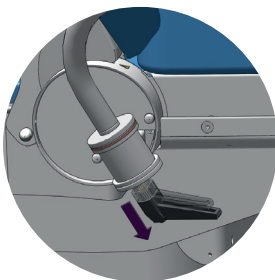
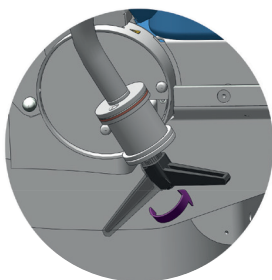
Adjustment of the distance between the electric footrests

(only on models 15630-01 and 15640-01)

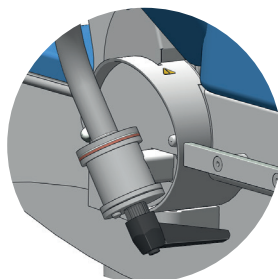
The distance between the footrests can be adjusted manually.



Indexing of the clamping handle



Resting position of the clamping handle



Safety note

Once the distance has been set, the clamping handle must be oriented as shown to prevent collisions.

Using the individual control

Protection at the electric footrests

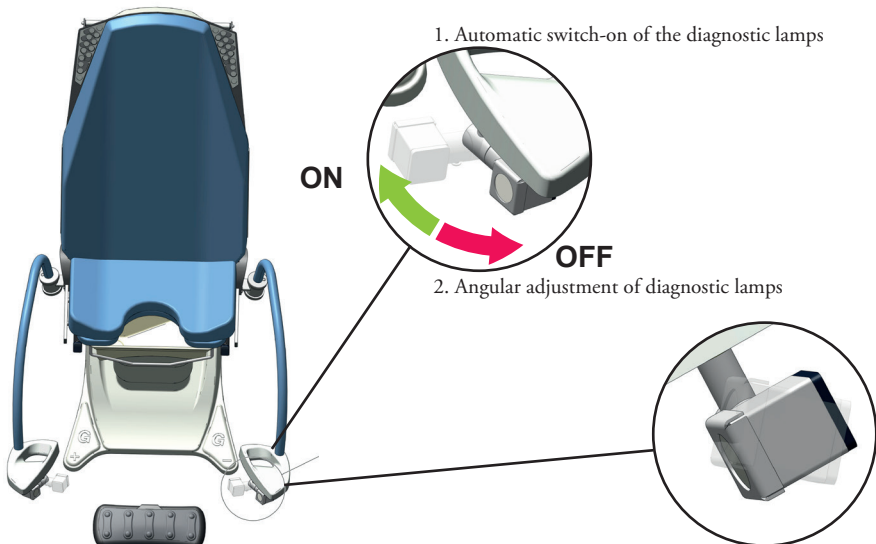
(only on models 15630-01 and 15640-01)

In order to avoid any risk of pinching between the footrests and the floor, the footrests are equipped with sensors that detect any objects that might be crushed. Therefore, when contact is made with one of these objects, all the actuators of the medical device stop, then the DM rises 5cm to unlock this object.



Use of integrated lamps

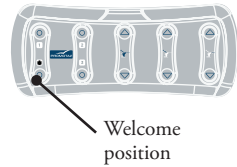
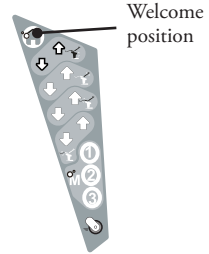
(only on model 15640-01)



Using the individual control

Welcome position

The position can be recalled by pressing once on the button .
NB: This position cannot be modified.




Placing on castors

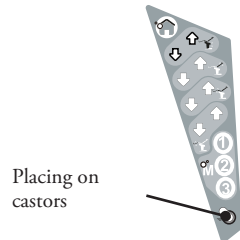
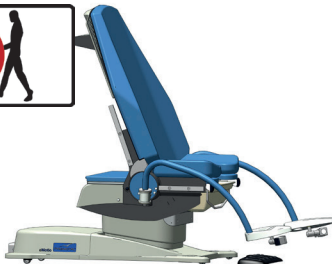
This position can be recalled by **pressing and holding** the button .

Return to a standard position by pressing once on the button .

NB: This position cannot be modified.



Safety note
Unplug the power supply cable before moving the MD.




After placing the device on castors, it may be moved by tipping it backward and manoeuvring it using the top of the backrest.

Never place castors on the device while a patient is seated on it.

Using the individual control

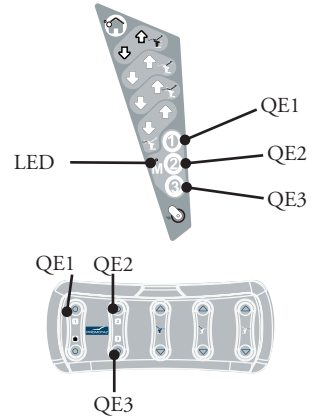
Position QE1, QE2, QE3

The keys QE1, QE2 and QE3 are factory-set. These are modifiable. To register a new position on the QE1 (or QE2 or QE3) button, it will move the DM to the desired position, then simultaneously press the button  and QE1 (or QE2 or QE3) for 6 seconds.

This will be registered when the LED flashes.

The recall is made by pressing the QE1, QE2 or QE3 key once.

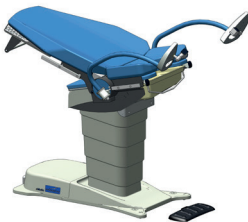
NB: This position cannot be modified. It is strictly prohibited to use the medical device to transfer patients.



QE1 (factory setting)



QE2 (factory setting)



QE3 (factory setting)



Using the individual control

Seat inclination control pedals

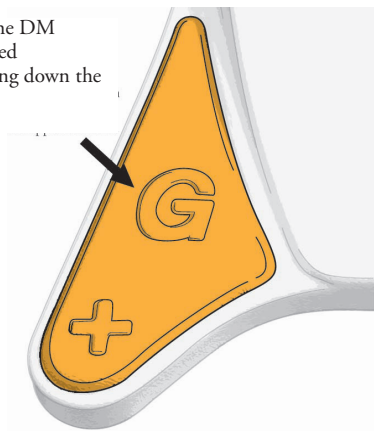
Using the pedals

The inclination is adjusted using the controls located on the edge of the housing.

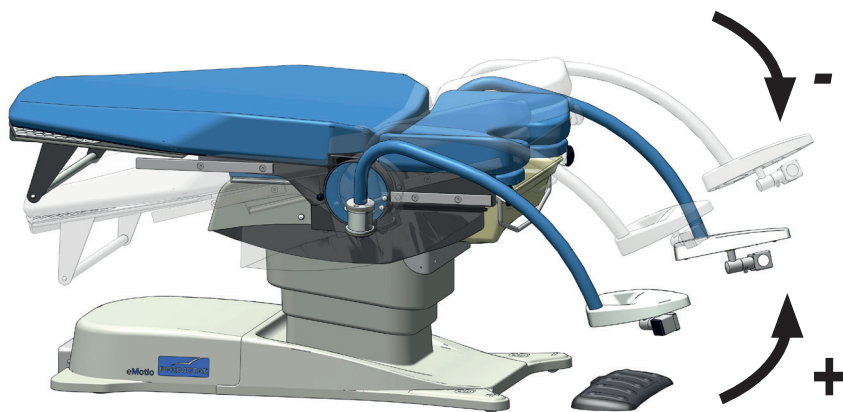
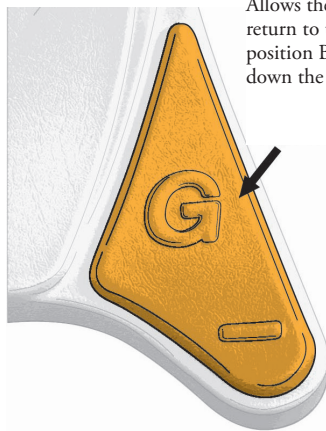
Press on the pedal **G+** to obtain an inclination.

The return to the horizontal position is done using the opposite pedal.

Allows the DM to be tilted
By holding down the button



Allows the DM to return to the horizontal position
By holding down the button



Safety note

For safety reasons, if the DM is in the down position, it will rise when the 20cm tilt is commanded and then tilt up

Using the gMotic

Paper roll

Installation of the paper roll holder

The paper roll is positioned on an axis with spring clips at each end. Press these springs to remove the paper roll.

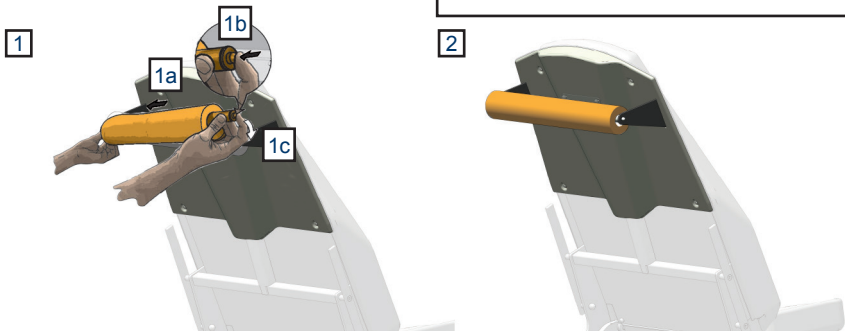
Cover the upholstery with paper before use.



Caution follow the indications provided well obtain a greater longevity of the material.



Caution Hold the paper roll holder during this intervention



Removing the upholstery

Raise the base by doing the 2 studs (P).

(Operation to be carried out for regular cleaning)

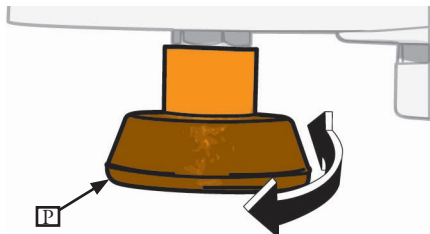


The levellers

Adjustment of the levellers

Only concerns the 2 rear levellers.

Screw or unscrew the leveller (P) to the required height.



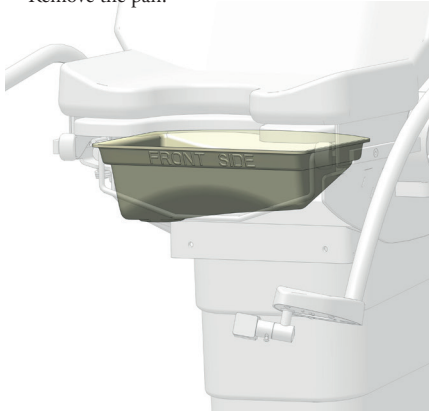
Using the gMotio


Plastic pan


(6 liters)

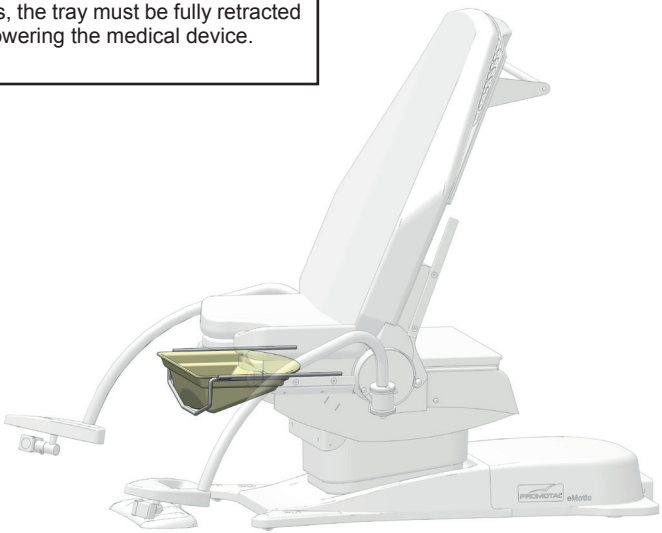
The plastic pan is removable to ensure easy cleaning, it can be moved towards the front and also be placed under the seat.


- Pull the handle underneath the seat, to move the pan forward.
- Remove the pan.



 Caution ensure the plastic pan is correctly positionde.

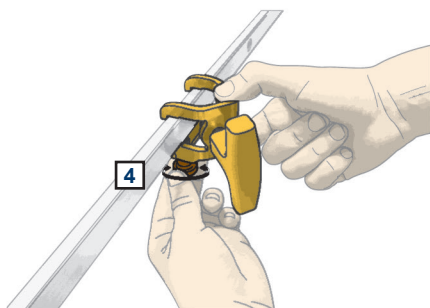
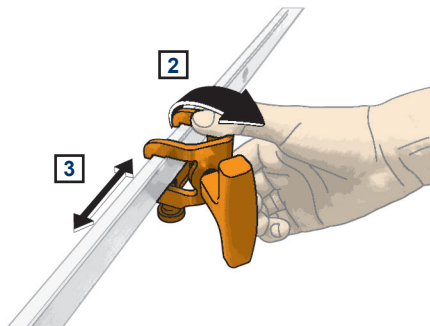
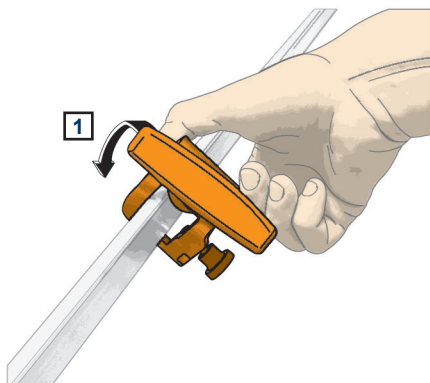
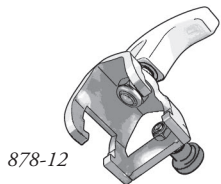
 **Safety note**
For safety reasons, in order to prevent collisions, the tray must be fully retracted before lowering the medical device.



 Only accessories designed and provided by Promotal for this medical device are authorised for use.

Sliding clamp for 25 x 10 rail (reference 878-12)

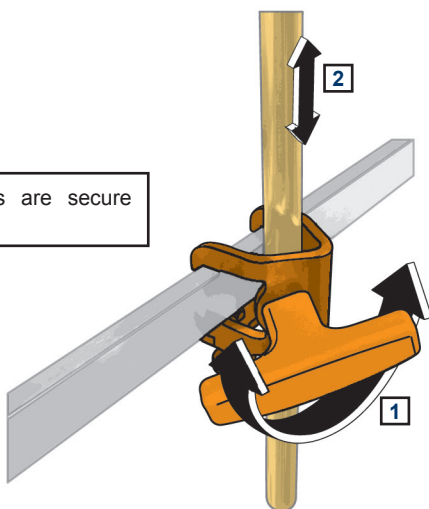
Installing the clamp



Using the clamp




Check that the parts are secure before use.




Pair of stirrups

(reference 850)

Installing the stirrups

 Ensures good clamping with each use

 **Caution**
Clamps are necessary to install these accessories.
(réf. : 878-12)


(For model 15620-01)




Pair of legrests GOEPEL

Installing the legrests

(For model 15620-01)

 Ensures good clamping with each use


 **Caution**
Clamps are necessary to install these accessories.
(réf. : 878-12)




Pair of legrests

Installing the legrests

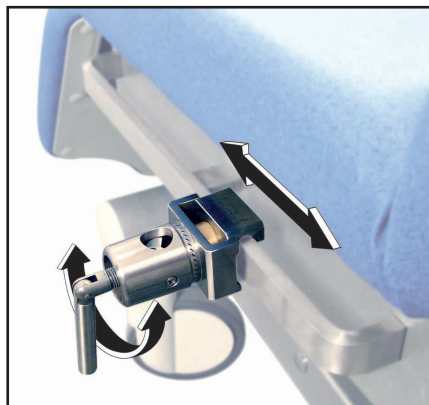
(For model 15620-01)

 Ensures good clamping with each use

 **Caution**
Clamps are necessary to install these accessories.
(réf. : 878-12)

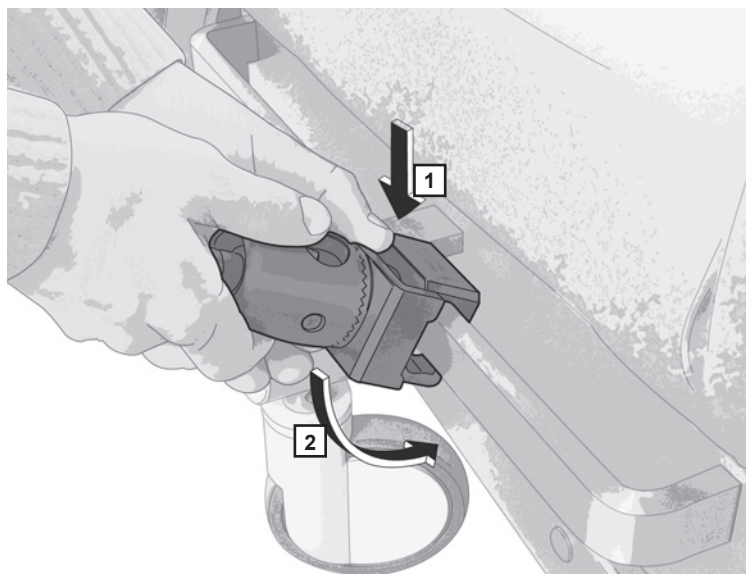



Rotating clamp for 25 x 10 rail (reference 879-10)




Installing the clamps

Insert the clamp on the rail.



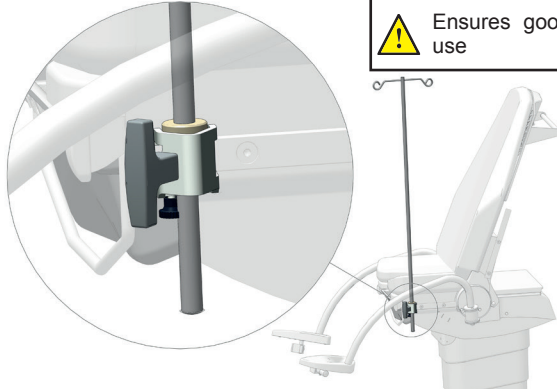
 Tighten the screws before use.


 Only accessories designed and provided by Promotal for this medical device are authorised for use.


2 hook IV pole

(reference 985-01)

Installing the IV pole



Caution
 Clamps are necessary to install these accessories.
(réf. : 878-12)

 Ensures good clamping with each use

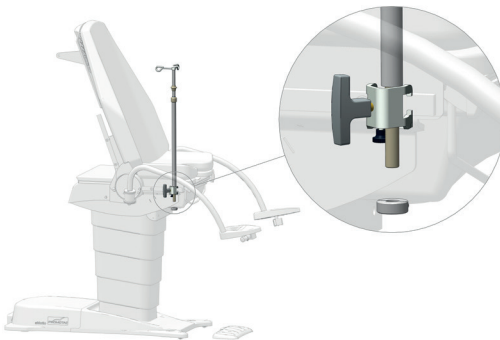
Auto-blocking I.V. pole


(reference 2985-01)


Installing the IV pole

Unscrew the screw handle.

Insert the I.V. pole and rescrew.



Caution
 Clamps are necessary to install these accessories.
(réf. : 878-12)

 Ensures good clamping with each use

Adjusting the height of the IV pole

The IV pole has 1 sliding stem.

Use the screw to adjust the height of the lower stem and the bolt to adjust the upper stem.



Manual control / 9 programmable positions

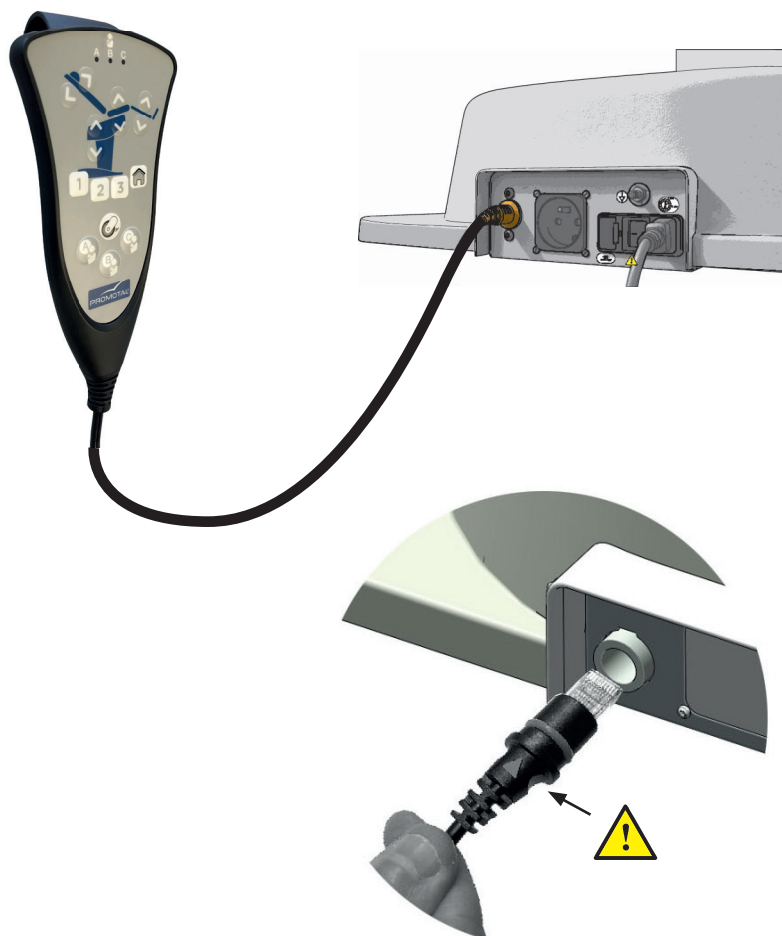
(reference 15031-10)



Safety note

When using the position memory function, movements can be interrupted by pushing on one of the command buttons for the DM or, as a last resort, by disconnecting the power supply cord.

Connection



Use

Adjusting the backrest angle (up)

Adjusting the height by holding the button down

Adjusting the backrest angle (down)

Adjusting the height by holding the button down

Adjusting the height (up)

Adjusting the height by holding the button down

Adjusting the height (down)

Adjusting the height by holding the button down

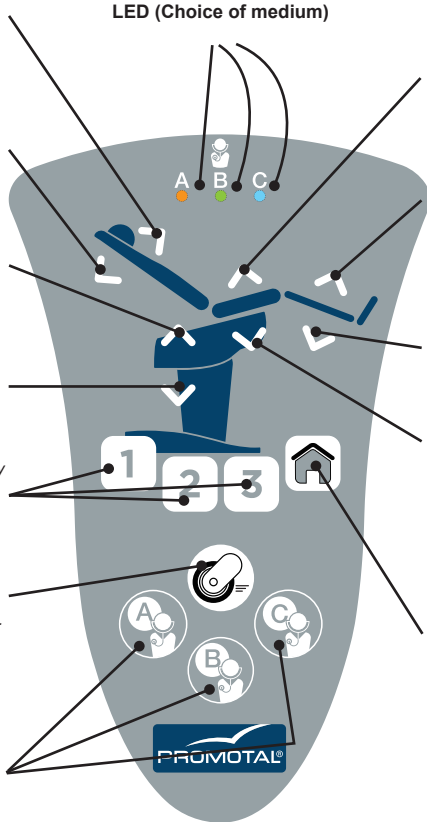
Positions QE1, QE2, QE3

This positions can be completely controlled by the user. At the touch of a button (push)

Wheel position

The position of the DM on the wheels allows it to be moved for ideal cleaning.

Choice of medium



LED (Choice of medium)

Adjusting the seat angle (up)

Adjusting the height by holding the button down

Adjustment of electric footrests (15630-01 & 15640-01)

Adjusting the height by holding the button down

Adjustment of electric footrests (15630-01 & 15640-01) (down)

Adjusting the height by holding the button down

Adjusting the seat angle (down)

Adjusting the height by holding the button down

Default position

Standard position for getting into and out of the DM (H: 57cm) at the touch of a button (push) NB: This position cannot be altered

QE = Quick Exam

Position QE1, QE2, QE3

The QE1, QE2 and QE3 buttons are factory-set but cannot be modified. To enter a new position for buttons QE1, QE2 and QE3, simply adjust the DM to the desired position and then press and hold the QE1 (or QE2, QE3) button for 6 seconds. This is stored once the LED located above the command bar blinks. The position can now be recalled by pressing once on the QE1 or QE2 or QE3 button.

Using the Doctor Choice Keys

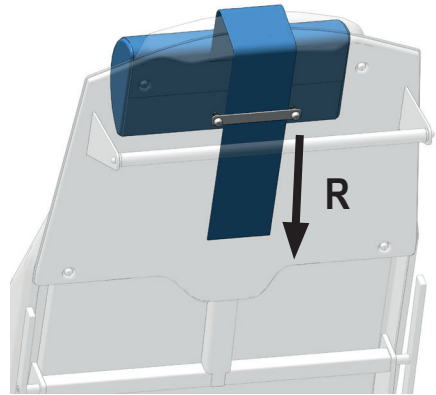
The first time the device is used, the choice selected by doctor A is active (orange LED). To switch to Doctor B (or C) simply press the B (or C) button for 7 seconds; the LED at the top of the hand control will change to GREEN (or BLUE if you have selected Doctor C). Once selection B (or C) is made, the QE1, QE2 and QE3 buttons will be automatically assigned to this doctor and you can memorize them by repeating the operations described above (USING Positions QE1, QE2 and QE3)

Head rest

(reference 11679-01)


Positioning

Push the head rest's band insert into the plate fixed to the back rest.



Adjustment

Just press on the head rest to lower it (D), or pull on the band insert to raise it (R).

 Only accessories designed and provided by Promotal for this medical device are authorised for use.

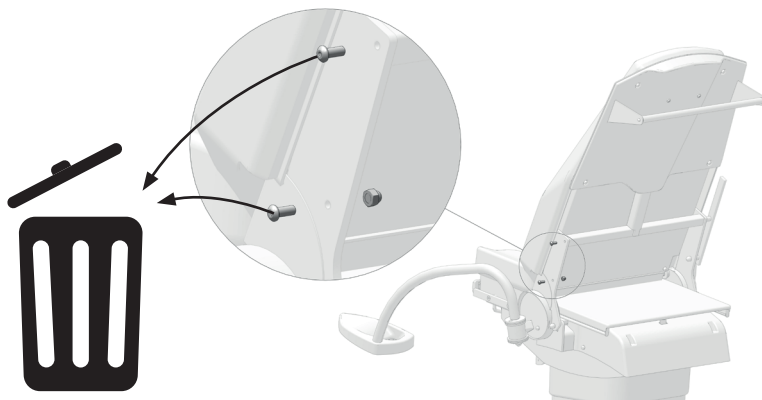
Pair of 25x10 stainless steel rails (Backrest) (reference 15879-01)

Removal of the screws present

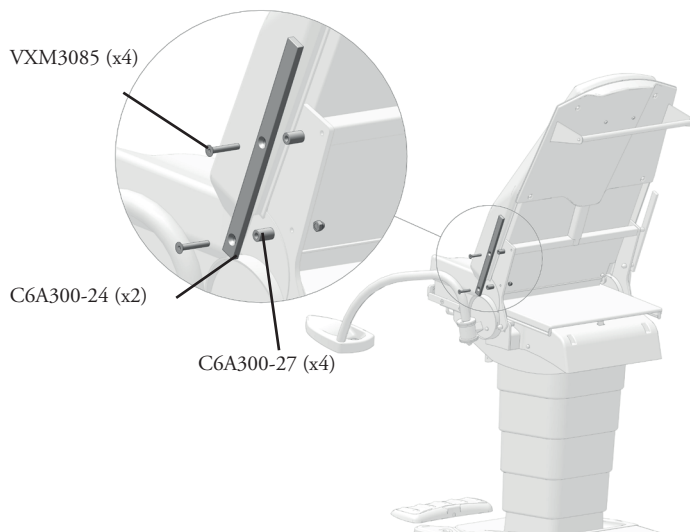
Box spanner of 13



N°5 Allen key

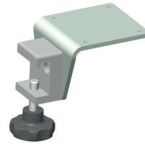



Assembly of the pair of rails

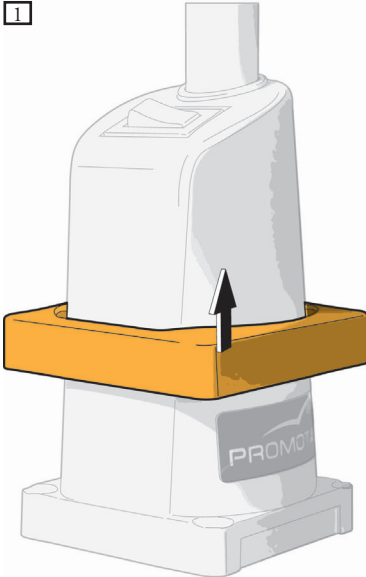


Lamp holder LID (reference 295-01)

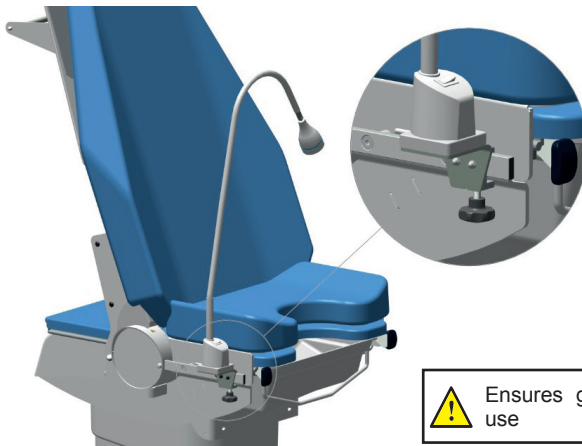
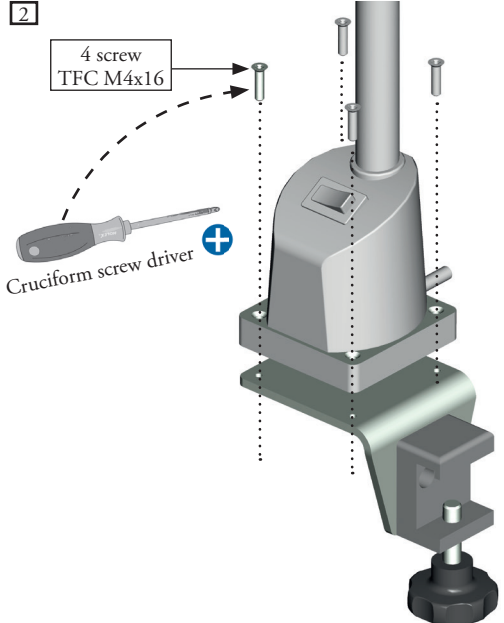
Fixing of the lamp on the support





 Ensures good clamping with each use



Installation on the table




 Ensures good clamping with each use

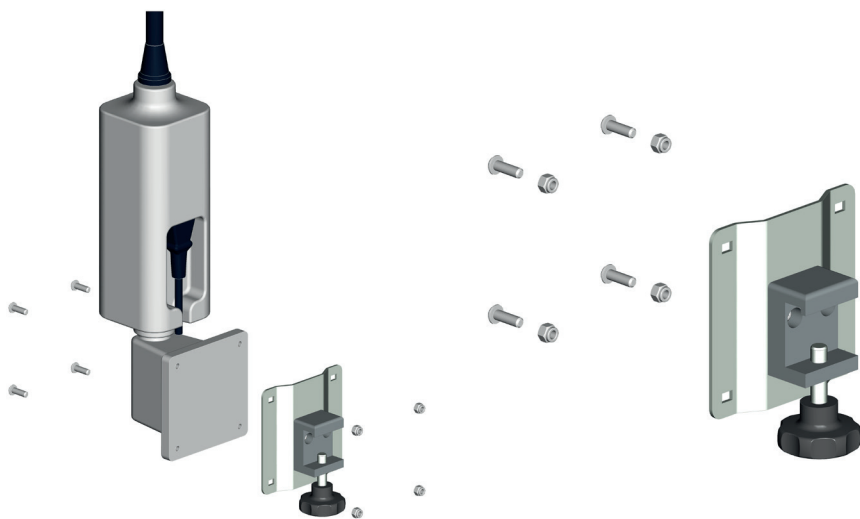
 Only accessories designed and provided by Promotal for this medical device are authorised for use.

Lamp holder Welch Allyn (reference 295-10)

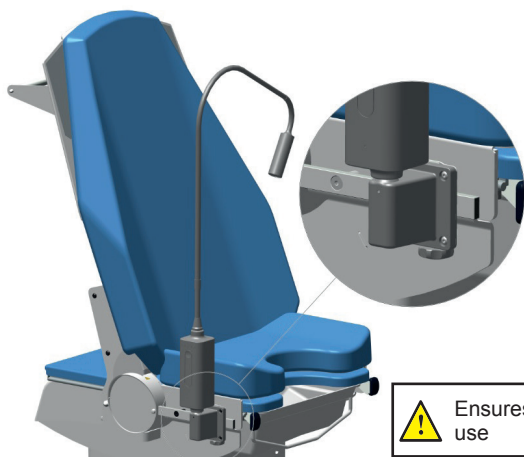
Fixing of the lamp on the support


Green Series™ Exam Light IV not included

 Ensures good clamping with each use



Installation on the table



 Ensures good clamping with each use

Leisegang Colposcope holder

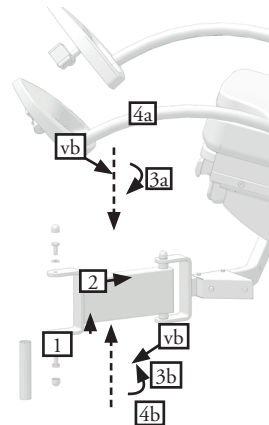
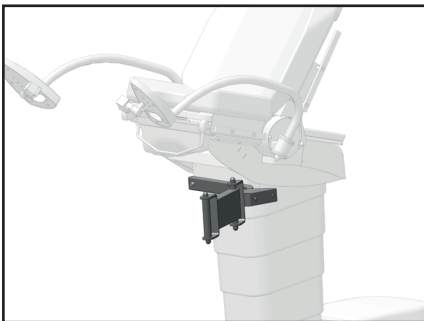
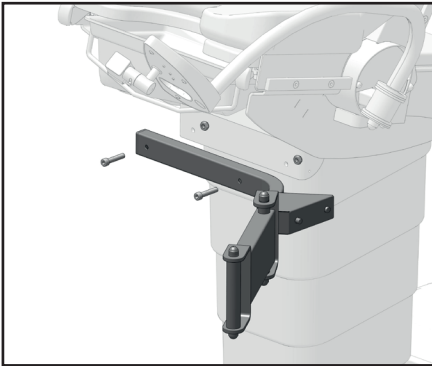
(reference 15810-01)

*Installing the colposcope holder
(Insert right or left)*




Incidence due to the adaptation of the colposcope support.

In order to avoid any risk of pinching between the colposcope and the DM, the impulse buttons QE1, QE2, QE3 and HOME will switch to maintained contact.



Tighten the 2 screws to the maximum (VB)


 Only accessories designed and provided by Promotal for this medical device are authorised for use.

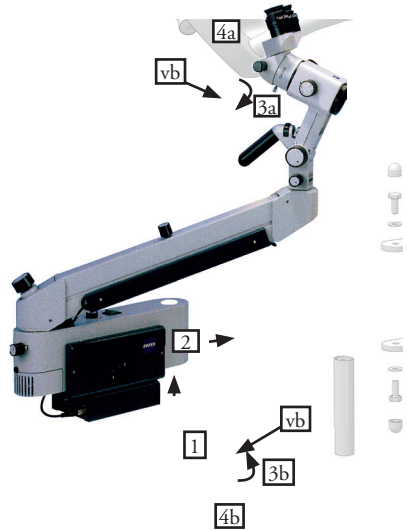
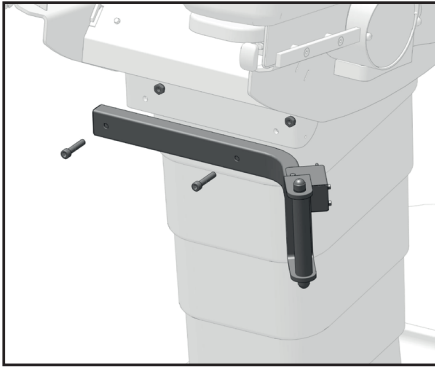
Zeiss Colposcope holder


(reference 15840-01)

*Installing the colposcope holder
(Insert right or left)*



 **Incidence due to the adaptation of the colposcope support.**
In order to avoid any risk of pinching between the colposcope and the DM, the impulse buttons QE1, QE2, QE3 and HOME will switch to maintained contact.



 Tighten the 2 screws to the maximum (VB)

Olympus Colposcope holder

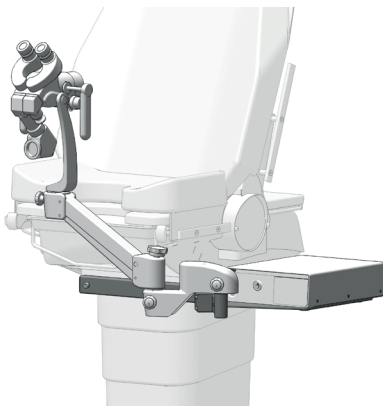
(reference 15850-01 right)

Installing the colposcope holder
(15851-01 left)

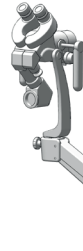


Incidence due to the adaptation of the colposcope support.

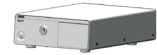
In order to avoid any risk of pinching between the colposcope and the DM, the impulse buttons QE1, QE2, QE3 and HOME will switch to maintained contact.



Supply «OLYMPUS»



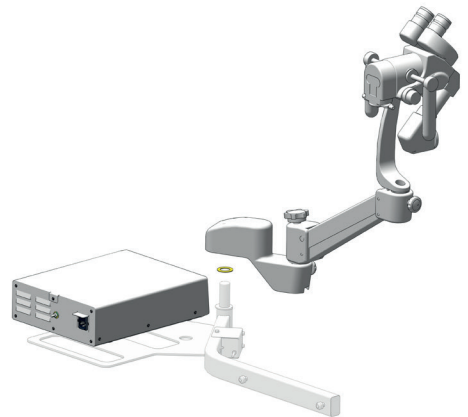
CCL-V1 Light Generator




OLYMPUS OCS-500



Tighten the 2 screws to the maximum (VB)



 Only accessories designed and provided by Promotal for this medical device are authorised for use.

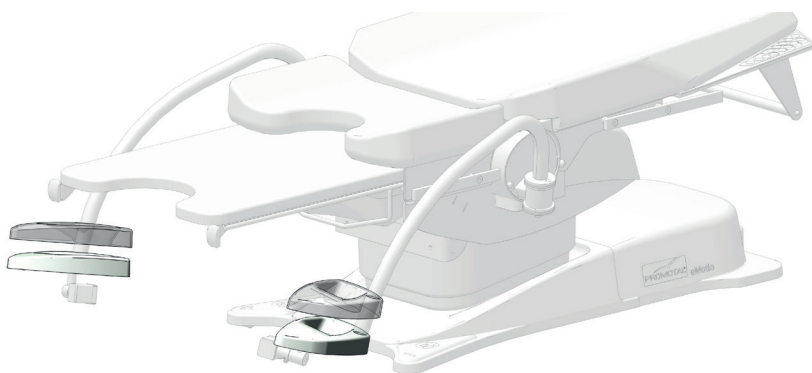
Pair of covers for legrests
(reference 861) for models 15626-01



Roll bar protection
(For models 15830-01 and 15840-01)
(reference 15861-01)
Protection reference



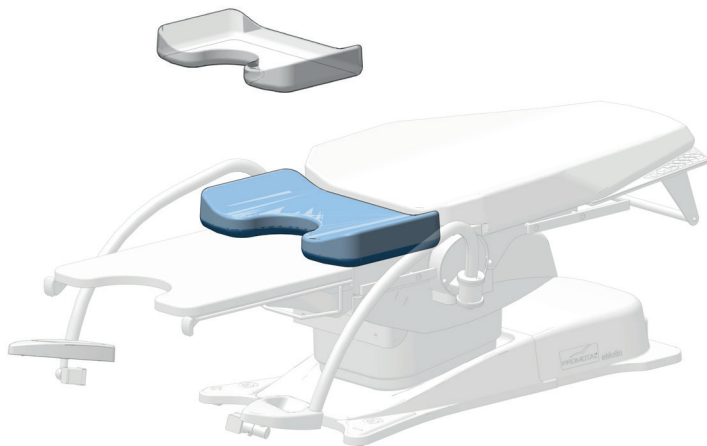
Foot Protection
(For models 15830-01 and 15840-01)
(référence 15905-01)
Protection reference



Seat protection

(reference 15906-01)

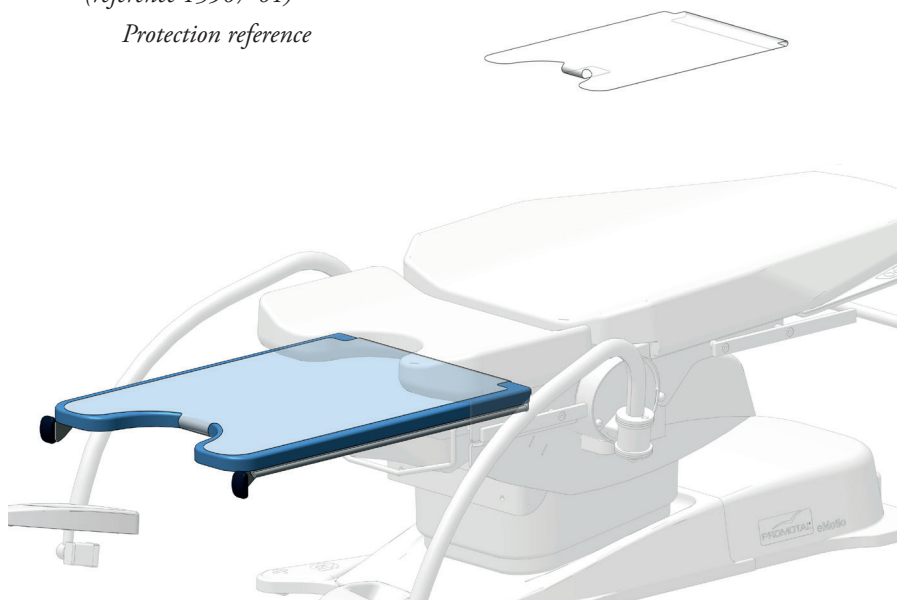
Protection reference



Leg rest extension protection

(reference 15907-01)

Protection reference



Options

Leg rest extension

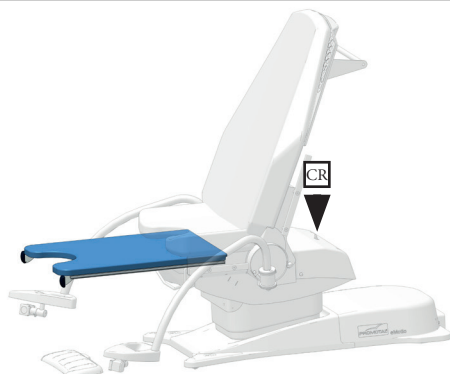
(reference 15672-01)

Using the leg rest extension



Safety note (only on models 15630-01 and 15640-01)

For safety reasons, in order to prevent collisions, the height adjustment of the electric footrests is blocked if the leg rest extension 15672-01 is not fully retracted.



To avoid any risk of pinching between the leg rest

and the floor, the retracted position of the leg rest is controlled by a CRposition sensor, if you press the keys that allow you to obtain the pre-programmed positions QE1, QE2, QE3 and HOME by impulsion, then the leg rest is not fully retracted, the movements will only be carried out with maintained contact, preceded by two audible beeps.

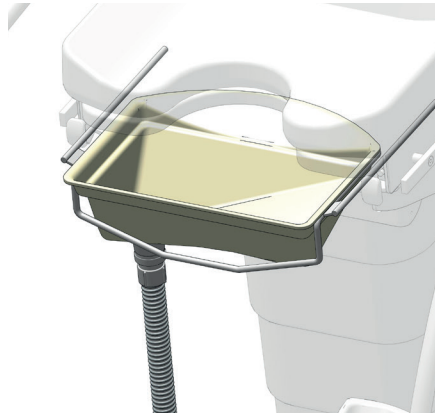
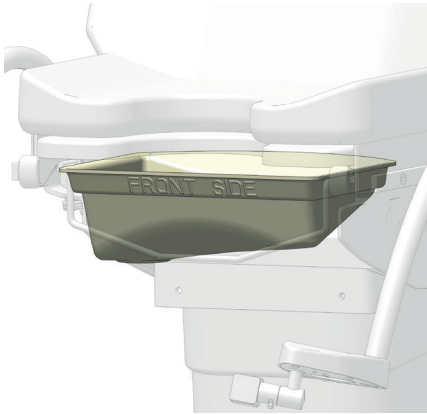
Plastic bowl with drainage tube

(reference 2061-30)

Sliding plastic removable bowl (6 liters) with drainage tube (150 cm).

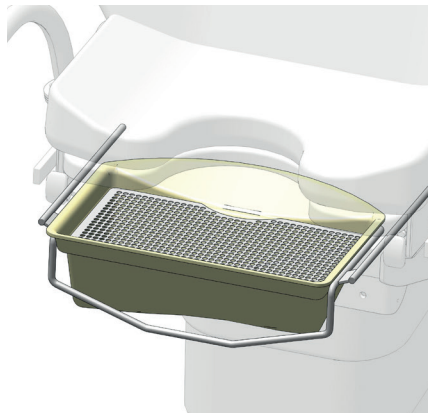
The plastic pan is removable to ensure easy cleaning, it can be moved towards the front and also be placed under the seat.

- Pull the handle underneath the seat, to move the pan forward.
- Remove the pan.



Splash guard cover

(reference 12677-01)



Options

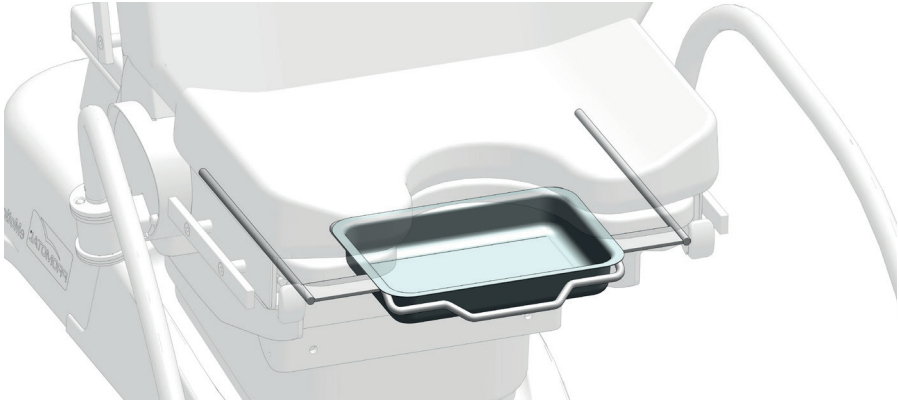
Stainless steel tray

(reference 15966-01)

Stainless steel tray (300 x 210 x 50) on rail

The tray is removable for easy cleaning, it can be moved forwards and folded under the seat.

- Grasp the handle underneath the seat to move the tray forward.
- Remove the tray.



Safety note

For safety reasons, in order to prevent collisions, the tray must be fully retracted before lowering the medical device.



Additional mains plug and equipotential terminal

(reference 40400-1 (EU, UK, US) 40400-2 (DK,IT,AUS,CH))

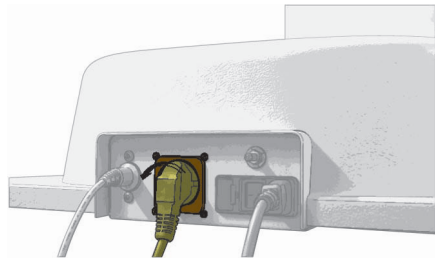
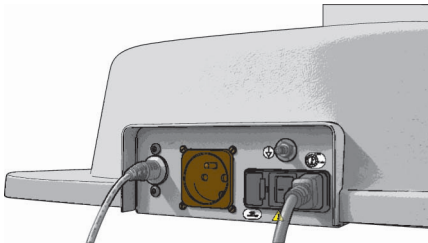


Safety note

By connecting a medical device to the additional mains plug, the unit becomes an Electro-medical system according to the standard EN 60601-1. The user must ensure that the EM system is in conformity with the standard EN 60601-1 (article 16).

Using the additional mains plug

- Characteristics : 120 V or 230 V depending on the country
2,5 A maxi



Lifespan of the Medical Device

This device is designed for a 10 year lifetime (except wear parts) under normal use conditions. This lifetime may vary according to frequency of use.

CHECK THE GENERAL CONDITION OF THE DEVICE AT LEAST ONCE A YEAR.

The wear parts are:

- The upholstery.
- The electric cylinders.

Promotal recommends replacing the wear parts after 5 years use maximum.

For all interventions, contact your usual dealer, indicating the Serial number of the device.



Material warning

Within the framework of maintenance, only the installation of components designed and provided for this MD by Promotal is authorised.

Compulsory / specific maintenance

Once a year, ensure that the following checks are carried out by a qualified technician (contact your dealer):

- Ensure that all screws are correctly tightened.
- Check the fixings of articulated parts.
- Ensure that the structure has not been deformed.
- Ensure the fuses are in good condition.
- Check that the power cable has not been cut or damaged.
- Check the different connections (excessive play, noise...)
- Note this information and the control date in the maintenance record.

Once a month :

- Check the correct operation of the safety devices (cf page 16).

Medical device end of service life

Your dealer is responsible for the recovery and end of life treatment of this device.

If necessary, do not hesitate to contact Promotal. We can propose solutions to treat this equipment in the best conditions.



Material warning

Upholstery whose coating is torn no longer provides an effective anti-bacterial barrier and must be replaced without delay.



Material warning

DO NOT dismantle THE DEVICE

If a fault is detected, immediately contact your dealer or the dealer's technical department (cf. Warranty Chapter) for a complete diagnosis. If you have a doubt, do not use the device.

Fuse replacement

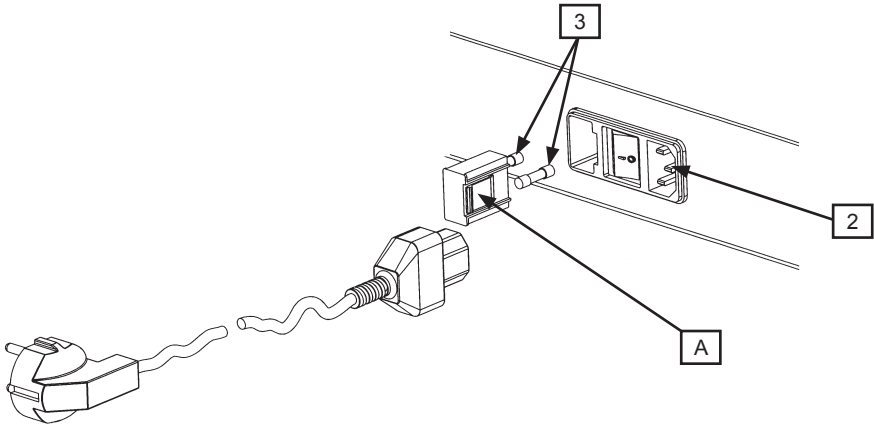
1. Remove all power to unit by unplugging unit's power cord.
2. Using a flat tipped screwdriver, gently release the plastic spring clip (A) on both sides of fuse drawer (1) then pull fuse drawer from fuse housing (2).
3. Pull both fuses (3) out of fuse drawer (1) and inspect. Check the fuses for any indication that they have blown ; i.e. burnt look, fuse cord melted through, etc. Discard fuses (3) if blown and replace them.



EQUIPMENT ALERT

Use fuses of the same voltage rating, amperage rating, and type. Failure to do so could result in damage to the equipment.

4. If necessary, obtain new fuse(s) (3). The replacement fuse(s) must be a 250 VAC, 6.3 amp, IEC 127 rated, 5 x 20 mm, Type T "Slo-Blo".
5. Insert fuse(s) (3) into fuse drawer (1).
6. Insert fuse drawer (1) into fuse housing (2) until fuse drawer snaps into place (both side of use drawer are locked into fuse housing).
7. Plug in power cord to equipment. If fuse blows again, call your Promotal distributor.



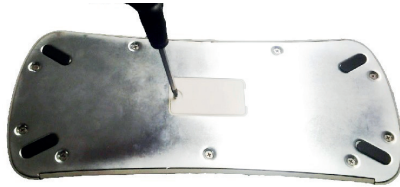
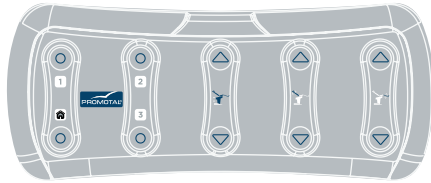
Replacing used batteries in the foot pedal

(battery reference (x2): LR03 - AAA Alkaline 1.5V)

To replace the batteries in this wireless foot pedal:

1. Turn the pedal over
2. Loosen the screw and remove the battery compartment cover
3. Remove the two used batteries and replace them with the two new batteries. (make sure to place them with their polarities correctly positioned.)
4. Replace the cover and tighten the screw.

Wireless foot pedal



MAINTENANCE NOTEBOOK

Product Reference :

Serial number (cf. label) :/...

Manufacturing date (start date of warranty) :

Date	Name of technician	Type of intervention - corrective action

If the product is returned to the retailer or manufacturer, please clean and disinfect the material.

Warranty Information

Warranty

Promotal will, at its discretion and expense, replace defective parts reported to Promotal within the applicable warranty period, and which, upon examination by Promotal, prove to be defective.

Obligations

Promotal will, at its discretion and expense, replace defective parts reported to Promotal within the applicable warranty period, and which, upon examination by Promotal, prove to be defective.

In accordance with CE regulation, Promotal's distributor is responsible for after sales service during and after the warranty period.

Exclusions

- This warranty does not extend to :

- (1) Spare parts and consumables.
- (2) Travel and labour expenses.
- (3) Breakdowns due to improper use, manifest neglect, or to moving the device.
- (4) Equipment whose original characteristics have been changed by the user.
- (5) Control units, hydraulic and electric jacks if opened by the user (seals broken).
- (6) Damage, breakdowns, failures, or defects attributable to outside causes (lightning, electric surges, floods, natural catastrophes, impacts, etc.) or to the presence of foreign objects.
- (7) Damage caused by improper hook-up or by the power supply, damage caused by corrosion or by gradual deterioration of the product.
- (8) Indirect damage related to loss of use and penalties generated by poor performance.
- (9) Aesthetic damage incurred by the outside parts of the equipment, which does not hinder proper operation, such as scratches, chips, and scrapes.
- (10) Devices whose serial number has been rendered illegible or has been changed or removed

Exclusive Remedy

Promotal's only obligation under this warranty is the replacement of defective parts. Promotal shall not be liable for any direct, indirect, special, incidental, exemplary, or consequential damages or delay, including, but not limited to, damages for loss of profits or loss of use.

No Authorization

No person or firm is authorized to create for Promotal any other obligation or liability in connection with the products.

THIS WARRANTY IS PROMOTAL'S ONLY WARRANTY AND IS IN LIEU OF ALL OTHER WARRANTIES, EXPRESS OR IMPLIED.

¹ Upholstery sets are warranted 1 year from manufacturing defaults.

EU Declaration of Conformity

We,

Promotal

22 rue de Saint Denis de Gastines

53500 Ernée – France

SRN: FR-MF-000001666

declare, under our sole responsibility, that the following electrical medical device:

Commercial name: gMotio 15620-01 / 15624-01 / 15626-01 / 15630-01 / 15640-01

Description: Electric Gynecological Table

Basic UDI-DI: 37014094GMOTIOGF

is a Class I medical device,

complies with the requirements of **Regulation (EU) 2017/745** of the European Parliament and of the Council of 5 April 2017,

complies with the following European directives:

- **Directive 2011/65/EU (RoHS 2)** of the European Parliament and of the Council of 8 June 2011
- **Delegated Directive (EU) 2015/863** of the Commission of 31 March 2015
- **Directive (EU) 2017/2102** of the European Parliament and of the Council of 15 November 2017

meets the applicable European standards,

bears, as such, the CE marking.

Year that the CE (Medical Devices) marking was initially affixed: 2020

Signed in Ernée,

On 26 May 2021

A handwritten signature in black ink, appearing to read "R. Mouradian", with a long horizontal flourish extending to the right.

Rudolf MOURADIAN

Chairman

Promotal

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