

LMErgo



User manual for LM-ErgoVet UltraLED



This manual is valid for:

LM-ErgoVet UltraLED 100741 VET LM-ErgoVet UltraLED 100751 VET

Manufacturer, marketing and sales

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1. INTRODUCTION

Foreword

This is User manual for LM-ErgoVet UltraLED product. Read and understand this manual before you use the product. If there are any questions regarding the contents of this manual, please contact LM-Instruments Oy. Keep User manual in place where it is easily available for all users.

General Requirements

Special precautions are necessary to this product regarding EMC (Electro Magnetic compatibility). It is necessary to install the product and put it into servicing referred to the EMC data in chapter "2.6. EMC Guidance and manufacturer's declaration" on page 7.

Portable and mobile RF (Radio Frequency) communications equipment can have an unwanted effect on the product.

Connect the product to electricity- and water supply which obeys the requirements in the Technical data.

Servicing

Only approved service persons are permitted to do servicing to this product. The maintenance procedures which are introduced in this manual are for the user. User must obey the maintenance measures to keep safe and efficient operation of the product.



2. SAFETY INSTRUCTIONS

2.1. Safety notices

The safety notices shown on this page are used to identify safety messages in these instructions:

WARNING	WARNING indicates a dangerous situation that, if not prevented, could cause death or injury.
	CAUTION indicates a dangerous situation that, if not prevented, could cause minor or moderate injury.
NOTICE!	NOTICE indicates a situation that, if not prevented, could result in material damage.



This general hazard symbol identifies important safety messages in this manual. Carefully read, understand and obey the messages. When you see this symbol, be alert: your and your patient safety is involved.



2.2. Electro magnetic compatibility

Special precautions are necessary to this product regarding EMC (Electro Magnetic compatibility). It is necessary to install the product and put it into servicing referred to the EMC data in chapter "2.6. EMC Guidance and manufacturer's declaration" on page 7. Portable and mobile RF (Radio Frequency) communications equipment can affect the product.

2.3. Safety considerations to install, servicing and to repair the LM-ErgoVet UltraLED unit

Connect the product to electricity- and water supply which obeys the requirements in the "8. Technical data" on page 28. Only persons with applicable professional knowledge are permitted to connect the unit to the water supply.

Only approved service persons are permitted to do servicing to this product.

In this manual are shown all the permitted maintenance procedures for the user, any other procedures are strictly prohibited. Before you do any maintenance work, read and understand maintenance instructions.

2.4. Intended and prohibited use

LM-ErgoVet UltraLED is designed for veterinary dentistry to removal of biofilm and calculus on teeth and other dental work where the ultrasonic vibration is beneficial. The unit must only be used by licensed veterinary dental professionals approved in the correct operation of scaling devices.

Do not use LM-ErgoVet UltraLED applications to human treatment or to other purposes where it is not intended. If you are unsure about your application, please contact your local dealer or place of purchase.



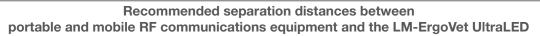
Do not make modifications to this product.

2.5. EMC Guidance and manufacturer's declaration

		r's declaration - electro			
÷		ne electromagnetic environme			
The customer or the user of	the LM-ErgoVet Ultra	LED should assure that it is u	used in such an environment.		
Emission test	Compliance	Electromage	netic environment - guidance		
RF emissions CISPR 11	Group 1	function. Therefore, it	raLED uses RF energy only for its interna s RF emissions are very low and are not erference in nearby electronic equipment.		
RF emissions CISPR 11	Class B	The LM-ErgoVet Ult	raLED is suitable for use in all		
Harmonic emissions IEC 61000-3-2	Not applicable	establishments, includ directly connected to	ding domestic establishments and those the public low-voltage power supply		
Voltage fluctuations/ flicker emissions IEC 61000-3-3	Not applicable	network that supplies	network that supplies buildings used for domestic purposes.		
Guidano	ce and manufacture	r's declaration - electro	magnetic immunity		
		ne electromagnetic environme nat it is used in such an enviro	ent specified below. The customer or the pomment.		
Immunity test	IEC 60601 test level	Compliance leve	I Electromagnetic environment - guidance		
Electrostatic discharge (ESD)	±6 kV contact	±6 kV contact	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity		
IEC 61000-4-2	±8 kV air	±8 kV air	should be at least 30%.		
Electrical fast transient/ burst	±2 kV for power suppl lines	y ±2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.		
IEC 61000-4-4	±1 kV for input/output	lines ±1 kV for input/output lines	ıt		
Surge	±1 kV differential mode	±1 kV differential mode	Mains power quality should be that of a typical commercial or hospital environment.		
IEC 61000-4-5	±2 kV common mode	±2 kV common mod			
Voltage dips, short interruptions and voltage variations on power supply input lines	<5 % U _T (>95 % dip in U _T) for 0.5 cycle 40 % U _T (60 % dip in U _T) for 5 cycles	<5 % U _T (>95 % dip in U _T) for 0.5 cycle 40 % U _T (60 % dip in U _T) for 5 cycles	Mains power quality should be that of a typical commercial or hospital environment. If the user tof the LM-ErgoVet UltraLED requires continued operation during power mains interruption, it is recommended		
IEC 61000-4-11	70 % U _τ (30 % dip in U _τ) for 25 cycles <5 % U _τ (>95 % dip in U _τ) for 5 sec	70 % U _T (30 % dip in U _T) for 25 cycles <5 % U _T (>95 % dip in U _T) for 5 sec	that the LM-ErgoVet UltraLED be powered from an uninterruptible power supply or battery.		
Power frequency (50/60 Hz) magnetic field	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercia or hospital environment.		



Guidance and manufacturer's declaration - electromagnetic immunity					
The LM-ErgoVet UltraLED is intended for use in the electromagnetic environment specified below.					
The customer or the user of the LM-ErgoVet UltraLED should assure that it is used in such an environment.					
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance		
			Portable and mobile RF communications equipment should be used no closer to any part of the LM-ErgoVet UltraLED including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.		
Conducted RF IEC 61000-4-6	3 Vrms 150 kHz to 80 MHz	3 V	Recommended separation distance: d = $1.2\sqrt{P}$		
Radiated RF	3 V/m	3 V/m	d = 1.2√P 80 MHz to 800 MHz		
IEC 61000-4-3	80 MHz to 2.5 GHz		d = 2.3√P 800 MHz to 2.5 GHZ		
			where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in metres (m).		
			Field strengths from fixed RF transmitters as determined by an electromagnetic site survey, ^a should be less than the compliance level in each frequency range ^b .		
			Interference may occur in the vicinity of equipment marked with the following symbol:		
NOTE 1 At 80 MHz and	800 MHz, the higher frequency	y range applies.			
NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.					
^a Field strengths from fixed transmitters, such as base stations from radio (cellular/cordless) telephones and land mobile radios amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the LM-ErgoVet UltraLED is used exceeds the applicable RF compliance level above, the LM-ErgoVet UltraLED should be observed to verify normal operation. If abnormal performance is observed additional measures may be necessary, such as reorienting or relocating the LM-ErgoVet UltraLED.					
Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.					



 $\mathsf{L}\mathsf{M}$

The LM-ErgoVet UltraLED is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. The customer or the user of the equipment can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the LM-ErgoVet UltraLED as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of	Separation distance according to frequency of transmitter				
transmitter	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2,5 GHz		
W	d = 1.2√P	d = 1.2√P	d = 2.3√P		
0.01	0.12	0.12	0.23		
0.1	0.38	0.38	0.73		
1	1.2	1.2	2.3		
10	3.8	3.8	7.3		
100	12	12	23		

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in metres (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE 1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

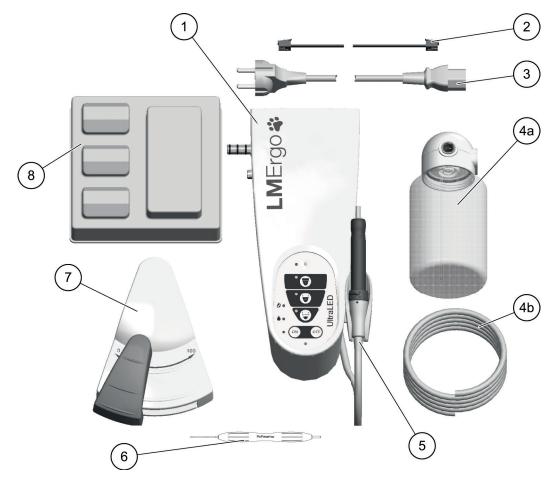
NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.



3. CONTENT OF DELIVERY

Carefully unpack your LM-ErgoVet UltraLED unit and verify all accessories and components are included to the content list below. If anything is missing contact your place of purchase.

3.1. Content of unit



1	LM-ErgoVet UltraLED unit	5	Scaler handpiece connected to unit
2	Foot control unit cable	6	LM-ProPower Fixer
3	Power cord	7	Foot control unit
4a (*)	500 ml medicament bottle	8	Scaler introkit (content varies depending on the order)
4b (*)	Water hose 6 mm (1/4")		

(*) depending of the type of the device

3.2. General description

LM-ErgoVet UltraLED is an effective piezoelectric scaler in one versatile and ergonomic appliance. The device's LED lights, advanced electronics, quality, and high-durability LM-DuraGradeMAX tips enhance the execution of procedures which require great precision. Ergonomically designed LM-ErgoGrip handpieces with soft silicone handles give the user a comfortable, relaxed grip as well as an excellent feel.



3.3. LM-ErgoVet UltraLED unit

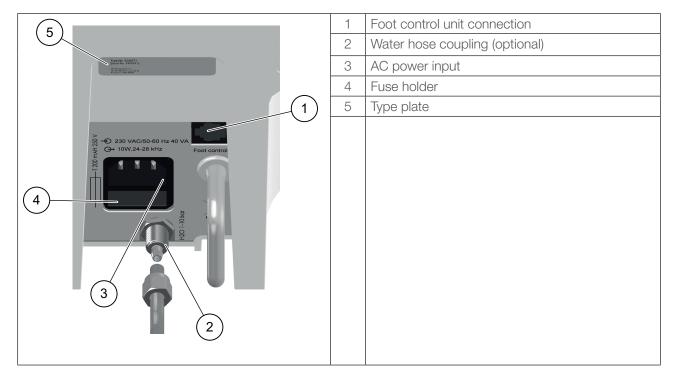


1	LM-ErgoVet UltraLED unit	5	Control panel
2	UltraLED handpiece (with an LM-ErgoGrip and a tip mounted)	6	Foot control unit
3	Water flow control ring	7	Depressurisation button (*)
4	Handpiece tubing	8	Medicament bottle (*)

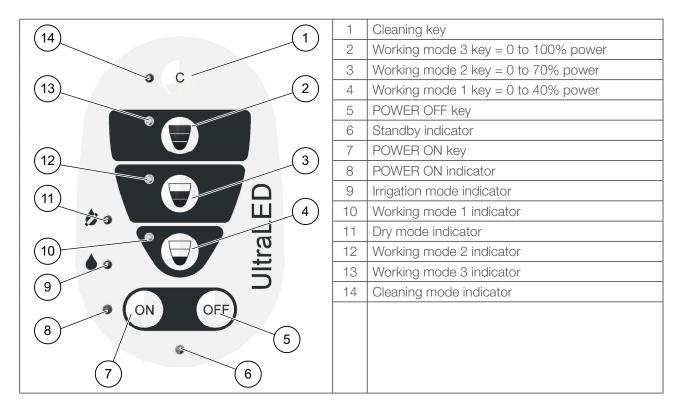
(*) not for tap water version



3.4. Coupling and type plate



3.5. Control panel



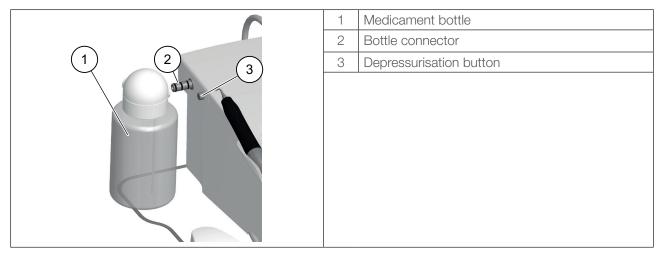


3.6. Medicament bottle (optional)

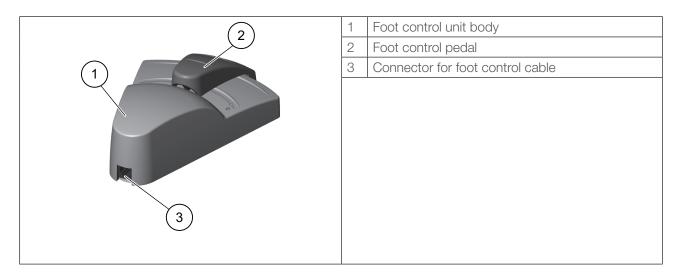
LM-ErgoVet UltraLED has a medicament dispenser system, making the device independent of a fixed water supply connection. The medicament bottle can be used for either medicament solutions or ordinary clean water. Approved medicament solutions:

- Clean water
- Cetylpyridinium chloride
- Chlorhexidine
- Essential oils
- Hydrogen peroxide, 3%
- Povidine Iodine, 10% solution
- Saline solution
- Sangurinara extract
- Sodium hypochlorite 1% solution

The unit contains an electrically driven air compressor. When operating, the compressed air forces the fluid from the bottle through the hose and to the handpiece and to the tip.



3.7. Foot Control unit





3.8. Symbols on the equipment

	Working mode 1
	Working mode 2
	Working mode 3
	Irrigation mode
	Dry mode
ON	Power ON
(OFF)	
	Power OFF
Scaler Weiter (M. Professor 10073) 751 British 10000 1 Scale (M. Scaler)	Example of type plate. The type plate is placed on the back side of the scaler unit.
CE	CE marking
	Caution
Ti	Consult accompanying documents
135°C	
135°C	Withstands autoclave temperature 135°C (275°F).
Τ	Type B applied part according to the degree of protection against electrical shock.
	Fuse
$ \rightarrow $	Input
\bigcirc	Output
X	Please do not throw the equipment into the domestic refuse. Please use the return and collection systems available in your country for the disposal of this product. The equipment can also be returned to the manufacturer for disposal.



4. TAKING LM-ERGOVET ULTRALED INTO USE

NOTICE!	The product should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the product must be observed to verify correct operation in the configuration in which it will be used.
NOTICE!	Special skills required. Consult an approved technician to connect the unit to the air and water supply.
NOTICE!	Do not put the unit on or adjacent to a heat source. Excessive heat can damage the electronics.
NOTICE!	Product is not suitable for use in the presence of flammable gases.

4.1. How to install the unit

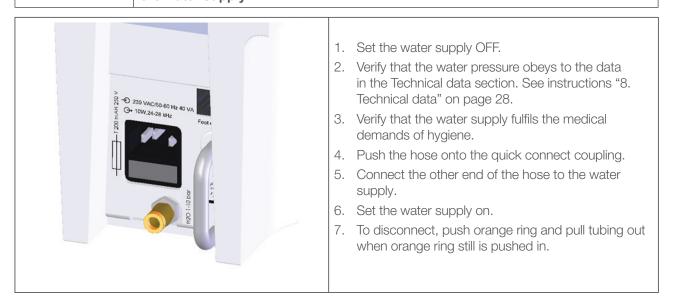




4.2. How to connect the unit for tap water (optional)

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NOTICE!
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Special skills required. Consult an approved technician to connect the unit to the water supply.



4.3. How to connect the unit to the supply current



ELECTRICAL HAZARD! Connect the unit to an AC power outlet supplied with a protective ground.

- 1. Verify that the voltage rating shown above type plate match the voltage of the AC power outlet.
- 2. Verify that the AC power supply is supplied with the protective ground.
- 3. Connect the power cable to the unit and the AC power supply. All indicator lamps illuminates for a short period when the unit does a self check.
- 4. The unit is standby when the green lamp is illuminated.

5. OPERATING INSTRUCTIONS LM-ERGOVET ULTRALED SCALER

LM

WARNING	PATIENT INJURY HAZARD! Do not use the scaler on patients with cardiac pacemakers. The scaler may disturb the function of the pacemaker.
WARNING	PATIENT INJURY HAZARD! New tips are not sterilized upon delivery. Sterilize tips before use. Non-sterilized tip may cause infection to the patient.
WARNING	ELECTRICAL HAZARD! Do not use damaged or worn-out handpiece. Damaged handpiece may expose the user or the patient for the electrical shock.
L	
WARNING	PATIENT INJURY HAZARD! Do not use a tip that is bent, altered worn more than 2 mm. Prolonged use may cause tip breakage and injury to the patient. Notice that ultrasonic instruments with small diameters are subject to breakage at any time. If not used correctly or with too much power or force the instrument will break

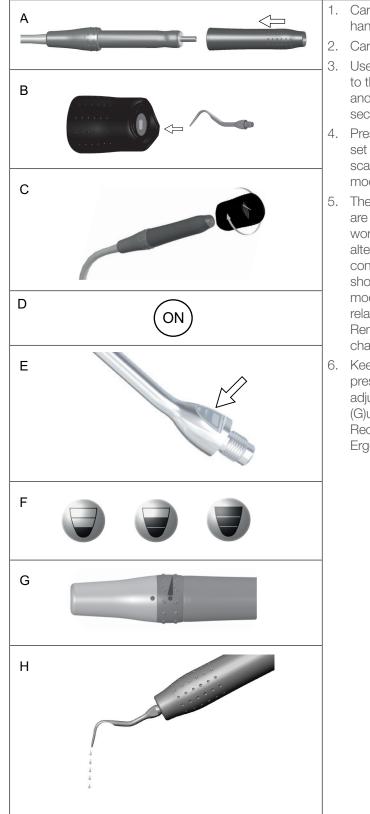
5.1. How to install LM-ErgoVet UltraLED scaler to use

PREPARATIONS (BOTTLE VERSION)

break.

- 1. Fill the medicament bottle with water or with approved medicament solution. Refer to the approved medicament solutions, see instructions "3.6. Medicament bottle (optional)" on page 13.
- 2. Turn the bottle cap onto the bottle and push it onto the connector.
- 3. Make sure that the unit power cable is connected.
- 4. Make sure that the unit is in standby mode, the green indicator lamp illuminates.

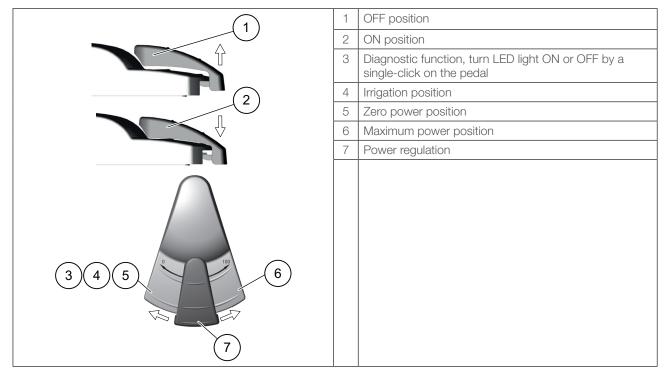




- 1. Carefully move the LM-ErgoGrip on to the scaler handpiece (A).
- 2. Carefully put the tip in the torque wrench (B).
- 3. Use the torque wrench and turn the tip clockwise on to the scaler handpiece (C). Tighten until resistance and the torque wrench slides. The torque wrench secures a correct tightening of the tip.
- 4. Press the key identified ON on the keyboard (D) to set ON the LM-ErgoVet UltraLED scaler. The blue scaling indicator lamp illuminates and the scaling mode is activated.
- 5. The indicators adjacent to the working modes are flashing to remind that you must select a working mode. If some blue indicators are flashing alternatively, make sure that the scaler handpiece is connected. If the problem stays, refer to the trouble shooting section. See the recommended working mode that is identified on the tip (E) and push the related working mode key (F) on the keyboard. Remember to choose right working mode if changing scaler tip during the treatment.
- 6. Keep the handpiece above the cuspidor bowl and press the foot control in the leftmost position and adjust the water flow with the ring on the handpiece (G)until the water is dripping from handpiece (H). Recommended flow: 20 ml/min, operating with LM-ErgoVet UltraLED scaler.



5.2. Foot control functions for scaler



5.3. Power and working modes for scaler

Scaling power

The power is controlled with the foot control from 0 to 100% in each working mode. See the recommended working mode that is identified on the tip. Press the related working mode key on the keyboard.

Dry mode for scaling

Press two times of the "ON" key to activate the dry mode for scaling without water/medicament.

Irrigation mode

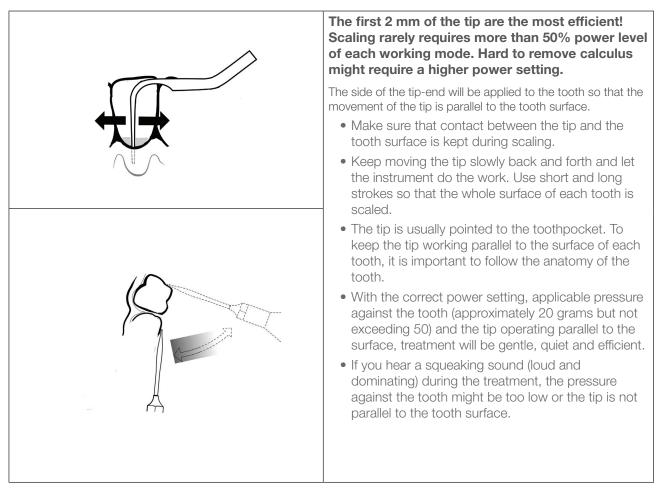
Press three times of the "ON" key to activate the irrigation mode. You can also activate the irrigation mode in the scaling mode by pressing down the foot control in the leftmost position.

When you press the "ON" key again and again, it will toggle between normal scaling, dry and irrigation mode.



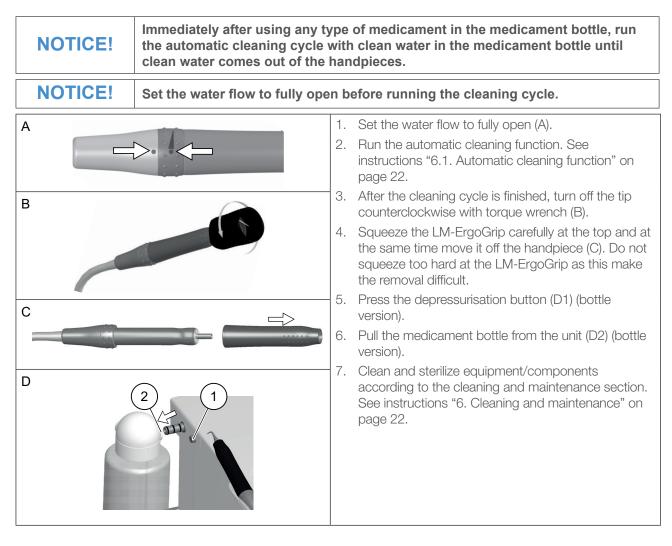
5.4. How to operate with LM-ErgoVet UltraLED scaler

	PATIENT INJURY HAZARD! Keep patients lips, cheeks and tongue out of the way of the activated tip. Tip contact could cause burns because of the friction heat.
NOTICE!	Do not use too much power when operating with ultrasonic instruments. If not used correctly or with too much force, instruments will break.
NOTICE!	Without cooling fluid the maximum operating time for the scaler handpiece is 2 minutes followed by a cooling-down period of 8 minutes. Operating without cooling fluid for more than 2 minutes could cause overheating of the scaler handpiece. After above cycle has been repeated 2 times, the scaler handpiece has to cool down for minimum 60 minutes.
NOTICE!	Increase the water flow if the handpiece feels too warm.
NOTICE!	Examine correct working mode from the tip.





5.5. After scaler treatment



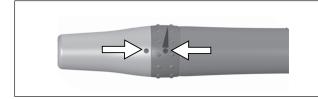
A short learning period is required by the user, as with any new technique, until the ideal angulations, soft tissue protection and an effective treatment can be achieved.



6. CLEANING AND MAINTENANCE

NOTICE!

Immediately after using any type of medicament in the medicament bottle, run the automatic cleaning cycle with clean water in the medicament bottle until clean water comes out of the handpieces.



Before cleaning and sterilizing, open water flow fully open of the scaler handpiece (see image).

6.1. Automatic cleaning function

- 1. Open the water control completely.
- 2. Put the scaler handpiece above the bowl.
- 3. Press the cleaning key "C" to start the cleaning cycle.
- 4. The cleaning cycle starts and stops automatically after 80 seconds.

6.2. How to clean the equipment and the components

NOTICE!	Autoclaving the handpiece regularly can decrease the life time of the scaler handpiece.
NOTICE!	Do not sterilize any scaler accessories using dry heat or chemical autoclaves. This may damage the material.

Clean off with a soft cloth and use a surface disinfectant suitable for hard plastics





Clean at max 65°C

Medicament bottle
Cap for the medicament bottle

Autoclave in steam at 134°C (max 135°C) for minimum 3 minutes

	Scaler tips
	Torque wrench
	LM-ErgoGrip Focus LED
LMErgo 🏶 🖁	LM-ErgoVet UltraLED scaler handpiece

6.3. Recommended cleaning procedure

Beginning of the day

Run the automatic cleaning cycle with clean water. See instructions "6.2. How to clean the equipment and the components" on page 22.

After each treatment

- Run the automatic cleaning cycle with clean water. See instructions "6.1. Automatic cleaning function" on page 22.
- Clean off the cover, control panel, handpiece and the hoses with a soft cloth. Use a surface disinfectant suitable for hard plastics.
- Clean the tip in an ultrasonic bath (40-50 °C), for minimum 3 minutes, before cleaning/sterilization. If ultrasonic cleaning is not possible, flush the tip with warm water.
- Clean the LM-ErgoGrip, the tip, and possibly the scaler handpiece and autoclave according to the clinic's routines.



End of the day

- Run the automatic cleaning cycle with clean water. See instructions "6.2. How to clean the equipment and the components" on page 22.
- Remove and clean medicament bottle and bottle cap at a maximum temperature of 65 °C.

Weekly (bottle version)

- Run the automatic cleaning cycle with an anti-microbial cleaning agent solution in the bottle. See instructions "6.2. How to clean the equipment and the components" on page 22. We recommend to use a separate bottle for the cleaning agent solution. Concerning exposure times of cleaning agent, follow instructions given by manufacturer.
- Before patient treatment. To flush the lines from cleaning agent solution, put clean water in the bottle and run the automatic cleaning cycle. Do automatic cleaning cycle again until clean water comes out of the handpiece.

6.4. Maintenance

NOTICE! A petroleum based lubricant on the O-rings can decrease their durability time.

Power cable

• Do an inspection of the power cable, cables and the handpiece hose daily to ensure that the equipment is in good condition without mechanical damage.

O-rings (bottle connector)

• Lubricate the O-rings regularly with water soluble lubricant.

Tips



PATIENT INJURY HAZARD! Do not use a tip that is bent, altered worn more than 2 mm. Prolonged use could cause tip breakage and injury to the patient. Notice that ultrasonic instruments with small diameters are subject to breakage at any time. If not used correctly or with too much power or force the instrument will break.

- Do an inspection of the tips. When a tip is bent, altered, or worn more than 2 mm it will loose performance and you must interchange the tip.
- Examine the tip length weekly by comparing the tip to a tip check card.

How to change fuses

4. Close the fuse holder.	3. Examine the fuses with ne Refer to the Technical dat	se holder on the rear side of the unit. fuses for damages. Replace damaged ew ones. Verify the fuse specifications. Technical data section. See section "8. ta" on page 28. se holder.
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7. TROUBLESHOOTING LM-ERGOVET ULTRALED

Type of problem	See chapter	
Unit does not give a result and no lights are lit on the control panel	7.1	
Lights are flashing on the control panel	7.2	
No tip vibration	7.3	
Weak tip vibration	7.4	
Insufficient or no water flow	7.5	
Tip does not fit smoothly onto the handpiece	7.6	
Difficult to remove the LM-ErgoGrip	7.7	
Handpiece is overheated	7.8	
Weak or no LED Light	7.9	

LM

7.1. Unit does not give a result and no lights are lit on the control panel

- 1. Make sure that the power cable is connected correctly and do a double check the voltage (120V/230V).
- 2. Examine the fuse and replace if necessary. See Maintenance-section of the user guide.
- 3. Examine that the wall outlet and the fuse panel of your clinic are OK.
- 4. If the problem stays contact your dealer for support and indicate Error Code E-X02.

7.2. Lights are flashing on the control panel

Error	Action
Four blue lights on the control panel are flashing at the same time.	Make sure that the foot control cable is connected at both ends and is not damaged.
The blue light adjacent to the ON-key is flashing alternately with three other blue lights on the control panel.	Make sure that the scaler handpiece is properly connected.
Three yellow lights are flashing in sequence.	The Working Mode has not been selected. Select the working mode by pressing one of the keys adjacent to the yellow lights.
The light adjacent to the C-key is flashing.	The cleaning function has been activated. Wait until the cleaning process has finished and the light stops flashing.

7.3. No tip vibration

- 1. Make sure that the tip is tightly tightened. Preferably use a torque wrench and tighten clockwise until the torque wrench slides and make sure that the tip is fully tightened. (The torque wrench secures a correct tightening of the tip.)
- 2. Examine that the green light below the keys on the control panel is lit. If it is not lit, please see Section 7.1.
- 3. Make sure that the device has been set on by pressing the ON-key and that the blue light adjacent to the ON-key is lit.
- 4. Make sure that the Working Mode has been selected by pressing one of the Working Mode keys and that one of the yellow lights on the control panel is lit.
- 5. If you have an additional handpiece, try replacing the handpiece.
- 6. If you have an additional foot control cable, try replacing the foot control cable.
- 7. If you have an additional foot control, try replacing the foot control.
- 8. If the problem stays contact your dealer for support and indicate Error Code E-S01.

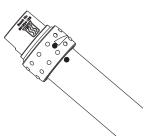


LM

- 1. Make sure that the tip is tightly tightened. We recommend that you use a torque wrench and tighten clockwise until the torque wrench slides. (The torque wrench prevents the tip from being overtightened.)
- 2. Examine that the tip is not worn out. Use a tip check card to evaluate wearing or try an unused tip. Use original LM tips for reliable performance.
- 3. If you have an additional handpiece, try replacing the handpiece.
- 4. If the problem stays contact your dealer for support and indicate Error Code E-S02.

7.5. Insufficient or no water flow when foot switch is activated

- 1. If the unit are equipped with a water bottle: Make sure that the medicament/water bottle is correctly connected i.e. the cap is tightened and pushed in all the way onto the connector. Examine that the O-rings on cap and connector are in good condition. Replace O-rings if worn. O-rings can be lubricated with a water soluble lubricant.
- 2. If the unit is connected with tap water: Make sure that the tap water hose is correctly connected to the back of the device and that the tap water system is OK (tap/valve is open and possible filter is OK).
- 3. Make sure that the Dry Mode is not activated i.e. that the blue light adjacent to the crossed-out water drop is not lit. If it is lit, press the ON-key.
- 4. Adjust the water control ring on the handpiece to fully open (dots aligned).



- 5. Try with different tip.
- 6. Use the LM-ProPower Fixer to examine that the water hose inside the handpiece has not stuck in the autoclave. Put the Fixer carefully into the handpiece from the hose connector side. (To prevent the hose getting stuck, the water adjustment on the handpiece should be set to fully open before sterilization in autoclave.)



7. If the problem stays – contact your dealer for support and indicate Error Code E-S04.

7.6. Tip does not fit smoothly onto the handpiece

- 1. Clean handpiece threads with compressed air and try with a new tip.
- 2. If the problem stays, the threads of the handpiece can be damaged and it is necessary to replace the handpiece. Contact your dealer and indicate Error Code E-S06.



7.7. Difficult to remove the LM-ErgoGrip from the handpiece

Hold the LM-ErgoGrip carefully near the lens and at the same time twist and move it off the handpiece.

7.8. Handpiece is overheated during use

- 1. Examine that the fluid flow is sufficient (minimum 20 ml/min).
- 2. If the problem stays, replace the handpiece and contact your dealer.

7.9. Weak or no LED light.

- 1. Examine that lens in the ErgoGrip is clear and clean. Clean or replace if necessary.
- 2. If you have an additional handpiece, try replacing the handpiece.
- 3. If the problem stays, contact your dealer for support and indicate Error Code E-S03.

8. TECHNICAL DATA

Manufacturer	LM-Instruments Oy, Norrbyn rantatie 8, FI-21600 Parainen, Finland
Model	LM-ErgoVet UltraLED
Classification	EN60601-1: Class 1, Type B
L×W×H	270 x 140 x 165 mm (without bottle)
Weight	2900 g
Voltage	100 Vac, 50-60 Hz
	115 Vac, 50-60 Hz
	230 Vac, 50-60 Hz
Primary fuse	T500 mA, 250 V, Ø5x20 mm (100 Vac)
	T400 mA, 250 V, Ø5x20 mm (115 Vac)
	T200 mA, 250 V, Ø5x20 mm (230 Vac)
Power cable	Separate with protective earth plug
Power consumption	Max. 40 VA
Scaler power consumption	Max. 24 VA
Scaler power output	Max. 10 W (24 kHz - 28 kHz automatic tuning)
Ambient temperature	Transport and storage -40°C to 70°C (-40°F to 158°F)
	Operation 10°C to 40°C (50°F to 104°F)
Relative humidity	Transport and storage 10% to 100%
	Operation 10% to 95%
Water supply pressure (version conn. to tap water)	1 - 10 bar (0.1–1.0 MPa, 14.5–145 PSI)
Water consumption	10 - 50 ml/min
Bottle volume (bottle version)	500 ml



9. WARRANTY

9.1. Warranty terms

The following warranty terms apply to the sale of LM-Instruments Oy's products (hereinafter "Products") to a purchasing company or individual by LM-Instruments Oy (hereinafter "Manufacturer").

Manufacturer here by warrants that the Products will be free from defects arising from faulty materials or workmanship for a period of twenty four (24) months from the date of purchase by a customer from Manufacturer's authorized dealer (hereinafter "Authorized Dealer"). The warranty period is exceptionally three (3) months for products with a life inherently shorter than 24 months due to normal wear and tear, for example ultrasonic scaler tips and tips of endodontics.

This warranty shall not apply to Products or parts thereof; which have been subject to abuse, misuse, negligence or accident or are not connected to proper power supply; to which any modification, alteration or attachment has been made without written consent of the Manufacturer; or which are installed or operated violating instructions for installation, use and maintenance; which are normally consumed in operation.

The sole and exclusive remedy under this warranty shall be limited to correct or circumvent the errors or to repair or replacement of defective parts of Products by the Manufacturer, EXW Manufacturer's factory, providing that a written claim of the defect is sent to the Manufacturer within the warranty period and the original part is returned to Manufacturer's factory by the Authorized Dealer, and Manufacturer's inspection establishes the existence of such a defect.

The customer must contact the Authorized Dealer from whom the products were purchased to request repair or replacement under this warranty and a written claim of the defects and send the original Product the Authorized Dealer.

This warranty is void if service or repair is performed by persons not authorized by the manufacturer.

Any Products not manufactured by the Manufacturer, carries only such warranty, if any, as given by any manufacturer thereof.

This warranty is the Manufacturer's only warranty in respect of the Products and the Manufacturer disclaims all other warranties, whether of merchantability, fitness for particular purpose or otherwise, guarantees and liabilities, express or implied, arising by law or otherwise. In no event shall the Manufacturer be liable for any general, consequential or incidental damages, loss of use or loss of profits by reason of the manufacturer's negligence or otherwise in connection with the sale, delivery, installation, repair or use of the Products.

The Manufacturer shall have no liability whatsoever to the Authorized Dealer or customer or any other person for or on account of any injury, loss or damage of any kind or nature, sustained by, or any damages assessed or asserted against, or any other liability incurred by or imposed upon the handling, use, operation, maintenance or repairs of Products by anyone other than the Manufacturer. This exclusion of liability does not apply pursuant to the laws on product liability in case of personal injury and property damage to privately used objects resulting from the Products.

9.2. Delivery Information

Device Model	LM-ErgoVet UltraLED		
Water connection	Bottle / Tap water connection		
Device Serial Number			
Purchase order number			
Order date			
Delivery date			

Installation information

Installation done (date)	
Installation done by (person)	
Installation location (address)	
Installation location (building and room number)	

9.3. Installation check-list

Task	Description	ОК	Initials	Date
Visual check	Delivery packing was undamaged			
Content of delivery	All the devices and part were according to the section "3. Content of delivery"			
Visual check	All the parts were intact			
Mains connection	Mains supply is in compliance with back panel label information (picture in section 3.4) voltage: V,Hz			
Water supply (bottle model)	Bottle filled with clean water, and bottle in place according to instructions in section 3.6			
Water supply (tap water model)	Tap water pressure is, and according to the specification 1-10 bar (0.1 – 1.0 MPa).			
	Water connection is close-knit and no indications of leakage is seen or heard			
Foot control	Cable connected to the main unit			
Scaler handpiece	ErgoGrip and tip assembled according to the instructions in "4. Taking LM-ErgoVet UltraLED into use".			

Instrument assembly is now ready for the operational test.



9.4. Operational test

Task	Description	Pass/fail	Initials	Date
Connect cable to mains	Green standby light is illuminated			
Turn the device on	Press "ON" on the control panel. Power on indicator is illuminated			
Test scaling with water -mode	Select 40% mode and test the foot pedal response. Use artificial denture in this test.			
Test scaling with water -mode	Select 70% mode and test the foot pedal response. Use artificial denture in this test.			
Test Scaler Irrigation mode	Select irrigation mode by pressing "ON". Test the water regulation of the handpiece by selecting irrigation mode and testing handpiece water flow ring positions. Continuous mist and dripping should be reached according the control ring angle.			
Test dry mode	Select 40% mode and test the foot pedal response. Use artificial denture in this test.			
Test dry mode	Select 70% mode and test the foot pedal response. Use artificial denture in this test.			
Test dry mode	Select 100% mode and test the foot pedal response. Use artificial denture in this test.			
Test diagnostic function	Turn LED on and off with the foot pedal			
Turn Device off	Press "OFF" on the control panel. Only standby LED is illuminated.			

Additional equipment needed: Artificial denture, cup for water flow, timer.

After all the above mentioned tests are passed the instrument is ready for the routine operation.

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