

PENTALED 30E

PENTALED 30N

PENTALED 63N

USER AND MAINTENANCE MANUAL

SURGICAL LUMINAIRE

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Introduction

Please read this manual carefully and thoroughly before using the Product in order to protect the **“TECHNICAL SERVICE PERSONNEL”** and the **“OPERATOR”** from damage.

CE Marking

This device is a Class I medical device within the meaning of REGULATION (EU) 2017/745 concerning medical devices (Annex VIII) as amended and supplemented.

Compliance

The manufacturer declares that this Product is in conformity with Annex I (General Safety and Performance Requirements) of REGULATION (EU) 2017/745 as amended and supplemented by IEC 60601-1, IEC 60601-1-2, IEC 60601-2-41 and documents this conformity by affixing the CE marking.

Manual validity

This manual is valid for the following models:

- PENTALED 30E in ceiling, wall-mounted, mobile version and respective battery versions
- PENTALED 30N in ceiling, wall-mounted, mobile version and respective battery versions
- PENTALED 63N in ceiling versions and respective battery versions

Customer service

Customer service is at your disposal for clarification of the Product, its use, locating spare parts, for service and/or warranty issues, and for any other questions.

- RIMSA P. LONGONI SRL
- Via Monterosa, 18/20/22 - 20831 - Seregno (MB) - Italia
- Tel.: ++39 0362 325.709
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- E-mail: info@rimsa.it

If the device causes the death or serious deterioration of the health of the patient and/or user, contact the manufacturer and the competent state authority where the event occurred.

Copyright

No part of this manual may be reproduced or translated without the written consent of RIMSA.

Translations

The original language of this manual is ITALIAN. For any translations, the original language of the manual shall prevail.

1 Legend

PRODUCT

The EM (electro-medical) EQUIPMENT to which this manual refers is a **SURGICAL LUMINAIRE FOR SURGICAL USE**. For ease of description in this manual, this EM EQUIPMENT will be referred to as **"Product"**.

OPERATOR

Professional medical personnel (e.g. professional health personnel, experienced patient caregiver).

RESPONSIBLE ORGANISATION

Entity responsible for the use and maintenance of EM equipment or an EM system (e.g. a hospital, an individual physician or an inexperienced person). Preparation and competence are included in the use.

TECHNICAL SERVICE PERSONNEL

Personnel (individuals or entities responsible to the RESPONSIBLE ORGANISATION) performing installation, assembly, maintenance or repair of the Product. In certain circumstances, their safety in accessing hazardous parts depends in part on their knowledge and competence to take appropriate precautions.

By way of example, but not limited to, the following professionals are considered TECHNICAL SERVICE PERSONNEL:

- Building Engineer, Surveyor, Building Contractor duly registered in the professional register (for construction work)
- Electrical Engineer, Electrical Technician qualified to practise as an electrician (for electrical works)

2 General safety information

This manual is an integral part of the Product as required by REGULATION (EU) 2017/745 as amended and supplemented. Please read and keep this manual close to the Product.



Risk of explosion.

The Product is not suitable for use in the presence of flammable anaesthetic agents.

RIMSA accepts no liability for any damage to persons or property resulting from the INSTALLATION/MANAGEMENT of the Product by anyone other than TECHNICAL SERVICE PERSONNEL or from the use of the Product by anyone other than the OPERATOR.

The installation of the Product is at the total expense and care of the RESPONSIBLE ORGANISATION; no burden or responsibility relating to the installation and/or commissioning of the Product can therefore be traced back to and/or in any way attributed to RIMSA.

The construction work to prepare the slab or wall, for Product to be installed on the ceiling or wall respectively, and the electrical work to prepare the electrical system to power the Product must be carried out in a solid and safe manner according to the rules of the art by the TECHNICAL SERVICE PERSONNEL.



Risk of electric shock.

To avoid the risk of electric shock, the Product must only be connected to a power supply network with protective ground.



Risk of electric shock.

The electrical layout of the room must comply with IEC 60364-7-710 and any national standards. A main switch with fuse protection or thermal-magnetic circuit breaker must be installed to ensure that the Product is de-energised.

3 Importance of personal safety

3.1 Intended use

SURGICAL LUMINAIRE FOR SURGICAL USE

The Product is an ELECTROMEDICAL APPARATUS that emits a light beam that can be directed independently of other light beams to illuminate a surgical procedure.

SURGICAL LUMINAIRE SYSTEM

A combination of several SURGICAL LAMPS that is SAFE TO FAIL and intended to be used during surgery to support the treatment and diagnosis of the PATIENT.

The optical radiation emitted by this Product complies with the exposure limits for the reduction of photobiological risk according to IEC 60601-2-41.

FIELD OF WORK

The Product provides good light intensity between 70 and 140 cm from the operating field.

3.2 Safety conditions (side effects)

Optical security

This product emits possibly hazardous optical radiation. Do not stare at the light emitted from the surgical luminaire. Eye injury may occur.

- Do not direct the light source into the eyes of the patient and OPERATOR.
- When the use of the Product is limited to the face (e.g. maxillofacial surgery, cosmetic surgery, ENT), it is mandatory to cover the patient's eyes with appropriate protection.



Possibility of glare and injury.

Failure to do so may result in glare and damage to the retina.

Electromagnetic interference

To avoid any significant risk of mutual interference due to the presence of the Product during specific examinations or treatments, please refer to the EMC declaration section in the user and maintenance manual.

Misuse



Placing objects on the Product is prohibited.

- Do not place, hang and/or rest any objects on the Product. Failure to comply with this requirement may cause such objects to fall into the operating area.
- Do not hang on the Product with your weight. In the case of the mobile version, do not climb, hang, lean, push or lie on the Product. Failure to comply with this requirement may result in damage to the Product, nearby devices and personnel.
- Do not cover the Product's lighthouse during operation to avoid overheating.
- Prevent Product parts from colliding with each other or with other neighbouring equipment. A collision may cause plastic or paint parts to detach from the Product and fall into the operating field.

Undesirable effects due to overlapping light fields

Overlapping the light fields of several lamp heads could lead to a temperature increase in the operating field and thus to a risk of dehydration and/or tissue damage.



Possibility of dehydration and tissue damage.

In the case of reduced blood supply with the principle of tissue dehydration, reduce the light intensity.

3.3 Environmental conditions



Risk of explosion.

The Product is not suitable for use in the presence of flammable anaesthetic agents.

During operation:

- the ambient temperature must be between 10 °C and 40 °C;
- relative humidity must be between 30% and 75%;
- atmospheric pressure must be between 700 and 1060 hPa;
- the altitude must be less than or equal to 2000 m.

4 General Information

4.1 Qualification of employees

Qualification of personnel to carry out Product operations:

Installation

Qualified installer and/or technician.

Use

Professional medical staff.

Cleaning

Thoroughly trained medical and paramedical staff.

Ordinary maintenance

Qualified technician in possession of the technical-professional requirements.

Extraordinary maintenance

RIMSA or TECHNICAL SERVICE PERSONNEL.

Assistance

RIMSA or authorised dealer.

Disposal

Comply with current waste disposal regulations. This Product must not be disposed of in normal waste bins. To avoid risks to the environment and health resulting from the dispersion of pollutants into the environment, separate the various internal components (e.g. iron, aluminium, plastic and electrical material) and take them to the appropriate centres for proper recycling.

4.2 Target population and interactions

Target population

The intended use makes the Product suitable for any population without constraints of age, weight, health or medical conditions.

Patients may be awake or unconscious, under local or general anaesthesia.

The reference population can also consist of animals.

Patient interaction

An active patient can only touch the Product's lighthead and swinging/pantograph arm accidentally, whereas such contact is excluded in the case of an unconscious or incapacitated patient.

Interaction with OPERATOR

The OPERATOR necessarily touches the Product's handle and keyboard, and occasionally the frame.

4.3 Explanation of the graphic symbols used in this manual

To emphasise their importance, some safety precautions are highlighted using the graphic markers below. Safety measures must be observed during installation, use and maintenance of the Product.



General warning sign.



General mandatory behavioural signal.



Generic prohibition signal.

Follow safety precautions before using or servicing the Product. Strictly following safety precautions improves the ability to safely and properly use the Product and helps prevent improper maintenance that can be dangerous and cause damage. The safety measures are indicative but not exhaustive; the OPERATOR, the RESPONSIBLE ORGANISATION and the TECHNICAL SERVICE PERSONNEL must develop their own skills in order to improve and supplement them.

4.4 Graphic symbols used on the Product

Description of any graphic symbols on the Product:























CE marking proving Product conformity with REGULATION (EU) 2017/745 as amended and supplemented



UL Classified Mark – MEDICAL – GENERAL MEDICAL EQUIPMENT AS TO ELECTRICAL SHOCK, FIRE AND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH AAMI ES60601-1:2005/(R)2012 and A1:2012/(R)2012 and A2:2021, IEC 60601-2-41:2021, CAN/CSA-C22.2 No. 60601-1:14 (Reaffirmed 2022), CSA C22.2 NO. 60601-2-41:23. Control No. E500026



UL Recognized Mark

	Date of manufacture (month and year)
	Address manufacturer
	Fuses used in the Product
	Follow the instructions for use
	Medical device
	Model Reference
	Serial number
	Unique device identifier
	Device identifier
	Authorised national representative
	Disposal
	Protective ground
	Connection point for neutral conductor
	Connection point for line conductor
	Accessed
	Off
	Stand-by and switch-on
	No pushing, leaning or lying on Product
	No climbing on Product
	Only move the Product with the arm lowered

5 Warnings for the Product Manager

5.1 Staff competence obligation

The RESPONSIBLE ORGANISATION must instruct the OPERATOR in the use, cleaning and maintenance of the Product. The instructions must be provided in written form on the basis of this manual.

5.2 Warranty and liability

RIMSA assumes no liability for unreliable operation of the Product if:

- The installation, authorised modifications and/or repairs are not carried out by TECHNICAL SERVICE PERSONNEL.
- The Product is not used in accordance with its intended use and with the user and maintenance manual.
- The room is not fit for use as a sanitary facility.
- The room is not constructed in accordance with the laws and/or regulations in force.
- The electrical installation of the premises does not comply with the appropriate regulations.

5.3 Checks to be carried out on Product before use



Carry out the electrical verification of the Product.

Before putting into service, carry out the electrical tests and the requirements specified in IEC 62353.

Before each use, in order to ensure Product safety and proper functionality, the OPERATOR must:

- clean/disinfect the Product in accordance with the provisions laid down by the competent national commission;
- check that the light emitted is stable and of adequate intensity;
- check that the lighthouse holds its position correctly;
- check that the swinging/pantograph arm holds its position correctly;
- check that there are no pieces or fragments of paint that can detach and fall onto the operating field. If they are present, remove them manually;
- check that the light source's protective shields are not damaged.

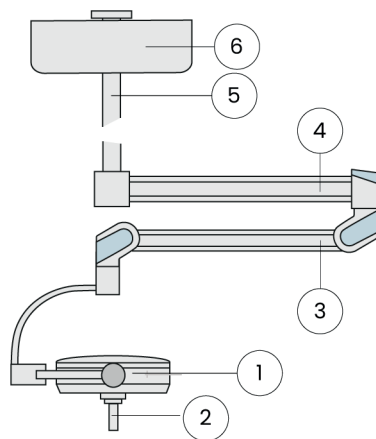
If one or more of these requirements are not met, please contact customer service.

6 Product Description and Operation

6.1 Product Description

6.2

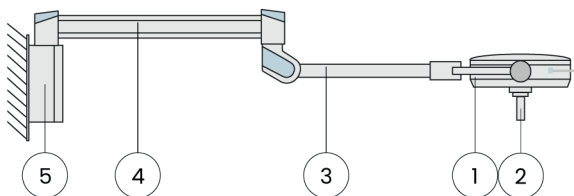
Ceiling version



- The Product is supplied inclusive of:
- lighthouse with single or double forks (1)
 - sterilisable handle (2)
 - swinging or pantograph arm (3)
 - horizontal arm structure (4)
 - tiges (5)
 - high or low cover with safety ring (6)
 - electrical panel.

Upon request, it is possible to equip the Product with a battery pack to allow operation even in the event of a power failure on the mains line. In this case, the electrical panel and battery pack are supplied together in a single box for each lighthouse belonging to the Product.

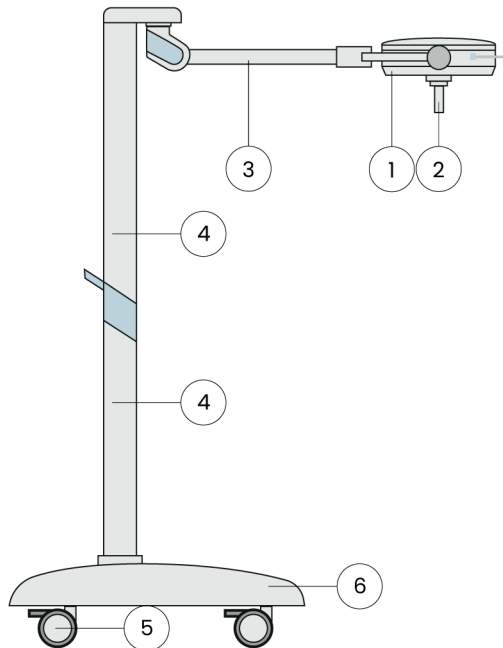
Wall-mounted version



- The Product is supplied inclusive of:
- lighthouse (1)
 - sterilisable handle (2)
 - swinging arm (3)
 - horizontal arm structure (4)
 - wall box (5)
 - electrical panel.

Upon request, it is possible to equip the Product with a battery pack to allow operation even in the event of a power failure on the mains line. In this case, the electrical panel and battery pack are supplied together in a single box for each lighthouse belonging to the Product.

Mobile version



The Product is supplied inclusive of:

- lighthouse (1)
- sterilisable handle (2)
- swinging arm (3)
- rods (4)
- base with wheels (5)
- base cover (6)
- electrical panel.

Upon request, it is possible to equip the Product with a battery pack to allow operation even in the event of a power failure on the mains line. For mobile Products, the Product with battery request must be made at the time of order as it is not possible to add the battery pack to pre-existing EM equipment.

Separable parts

- Sterilisable handle: refer to the section on cleaning, disinfection and sterilisation of the handle for assembly/disassembly instructions.

6.3 Description of Operation

6.3.1 Main switch

For ceiling versions, plan to place the circuit breaker in an accessible place and close to the Product, so that it can be switched off if necessary.

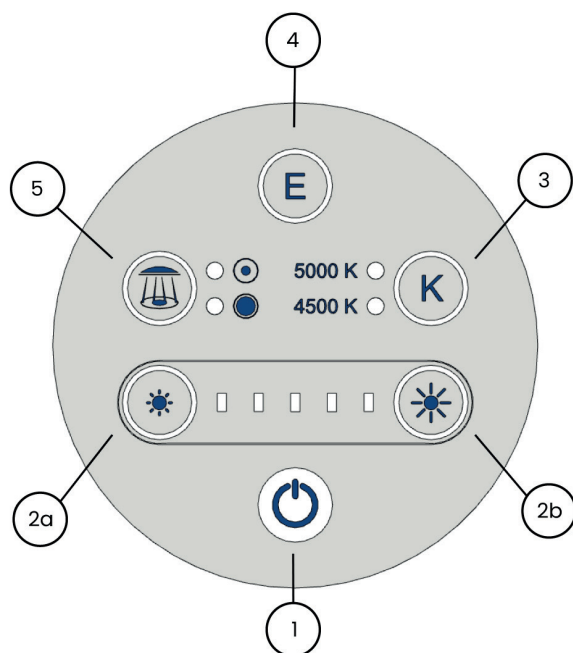
The wall-mounted and mobile version lamps are equipped with a green light switch for general on/off.



For wall-mounted and mobile versions, position the Product in such a way that it is easy to reach and disconnect the power plug in case of emergency.

6.3.2 Control keyboard SERIES E

The control keyboard for the functions is applied to the yoke of the Product.



By pressing the membrane keys, the following functions are activated:

- turning the Product on and off using the key (1);
- adjusting the light intensity using the keys (2). The level of intensity reached is displayed via 5 green microLEDs;
- adjusting the color temperature (CCT) using the key (3);
- activating the “Endoled” function using the key (4). This function can only be activated when the Product is off;
- adjusting the light diameter using the keys (5).

Illuminated area

The Product is designed to ensure a fixed light focus without the need for adjustment.

Synchronization/desynchronization of the Product

In the case of a Product with more than one lighthouse, it is possible to synchronize/desynchronize the lighthouses in terms of light intensity, color temperature, and/or light diameter.



The factory setting provides that the Product’s color temperature synchronization is active.

Light intensity synchronisation/desynchronisation

- to synchronise: with the lamp switched on and in surgical mode (not Endoled), set the light intensity to maximum; press and hold key (2b) until the relevant indicator LEDs flash.
- to desynchronise: with the lamp switched on and in surgical mode (not Endoled), set the light intensity to minimum; press and hold key (2a) until the relevant indicator LEDs flash.

Colour temperature synchronisation/desynchronisation

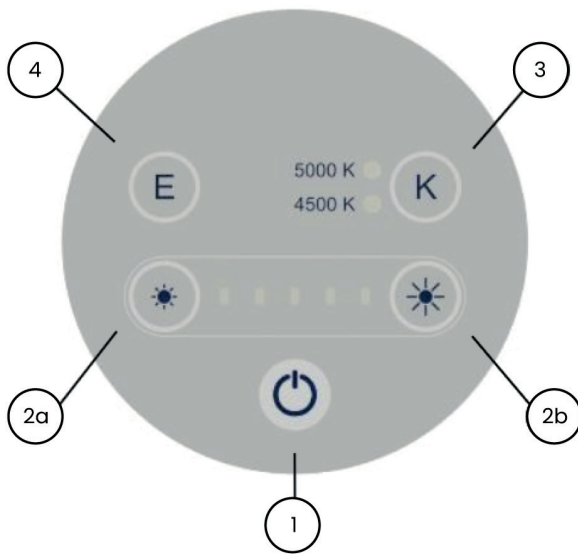
- to synchronise/desynchronise: with the lamp switched on and placed in surgical function (not Endoled), on the keyboard of a lighthouse of your choice press and hold the key (3) until the indicator LED flashes.

Light diameter synchronisation/desynchronisation

- to synchronise/desynchronise: with the lamp switched on and placed in surgical function (not Endoled), on the keyboard of a lighthouse of your choice press and hold the key (5) until the indicator LED flashes.

6.3.3 Control keyboard SERIES N

The control keyboard for the functions is applied to the yoke of the Product.



By pressing the membrane keys, the following functions are activated:

- turning the Product on and off using key (1);
- adjusting the light intensity using keys (2). The intensity level reached is displayed via 5 green microLEDs;
- adjusting the color temperature (CCT) using key (3);
- activating the “Endoed” function using key (4). This function can only be activated when the Product is off.

Upon turning on, the Product is by default set to surgical mode. Upon request, the Product PENTALED 63N can be set to turn on in endoscopic mode.

Light field adjustment

The mechanical adjustment of the light field is performed by rotating the sterilizable handle. By rotating clockwise or counterclockwise, the light field in the working area is widened or narrowed.

Synchronization/desynchronization of the Product

In the case of a Product with more than one lighthead, it is possible to synchronize/desynchronize the lightheads in terms of light intensity, color temperature, and/or light diameter.



The factory setting provides that the Product's color temperature synchronization is active.

Light intensity synchronisation/desynchronisation

- to synchronise: with the lamp switched on and in surgical mode (not Endoed), set the light intensity to maximum; press and hold key (2b) until the relevant indicator LEDs flash.
- to desynchronise: with the lamp switched on and in surgical mode (not Endoed), set the light intensity to minimum; press and hold key (2a) until the relevant indicator LEDs flash.

Colour temperature synchronisation/desynchronisation

- to synchronise/desynchronise: with the lamp switched on and placed in surgical function (not Endoed), on the keyboard of a lighthead of your choice press and hold the key (3) until the indicator LED flashes.

6.4 Product Handling

6.4.1 Movement mode

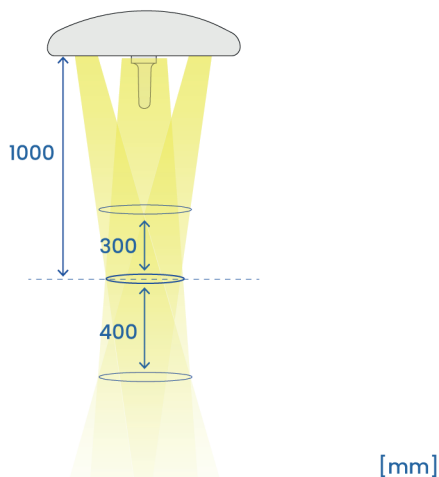
The Product can be moved using the sterilizable handle.



It is also possible to move the Product using the side handles.



6.4.2 Recommended working distance



The Product provides good light intensity at a distance of 70 cm to 140 cm from the operating field.

However, for the optimisation of light intensity, we recommend using the Product at a distance of 1 m.

6.4.3 Mobile version: handling recommendations

When it is deemed necessary to move the column, make sure to position the swinging arm downwards. If this warning is not heeded, the Product may tip over.



Possibility of Product tipping.

When handling the Product, it is recommended that the power cable be wound onto the cable reel on the vertical stem.

When positioning and using the Product, ensure that the power cable does not present a tripping hazard to the OPERATOR.



Possibility of tripping.



The mobile version is equipped with 4 wheels with foot brake to be operated to lock the Product in the desired position.

Press the brake pedal with your foot, without applying excessive force.



To deactivate the brake, lift the pedal with your foot.



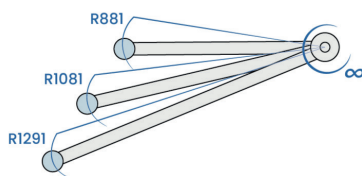
Possibility of damaging the pedal.

Do not strike or depress the brake pedal insistently once it has come to a standstill.

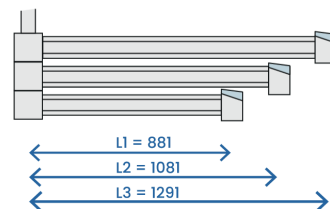
6.4.4 Dimensions and handling Product

Dimensions and handling horizontal arm structure

The dimensions of the Product depend on its configuration. The Product may have up to three lighthouse which will be arranged on the first, second and third horizontal arm of the structure respectively. In the following pictures, the length of the horizontal arm, measured in millimetres, is generically indicated with "L".



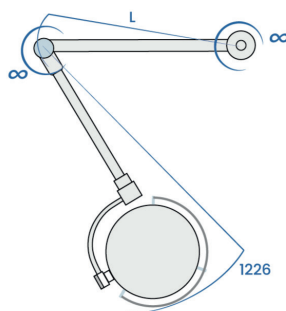
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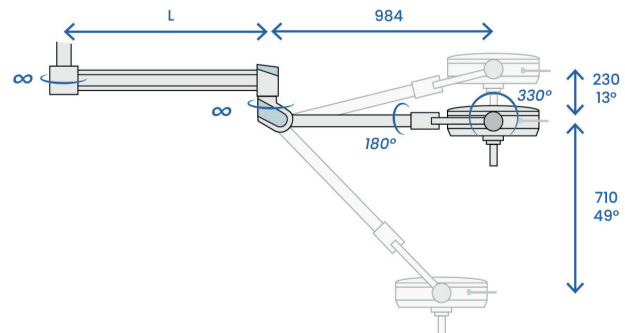
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Movement PENTALED 30E and PENTALED 30N

Ceiling version with single yoke

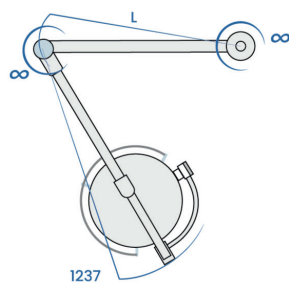


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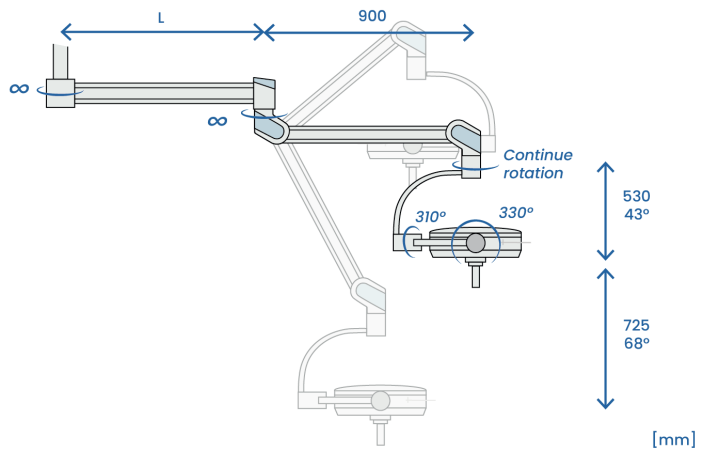


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Ceiling version with double yoke

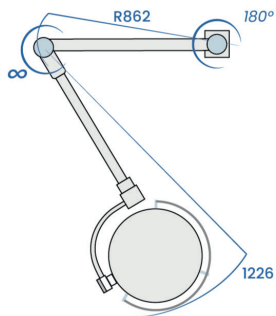


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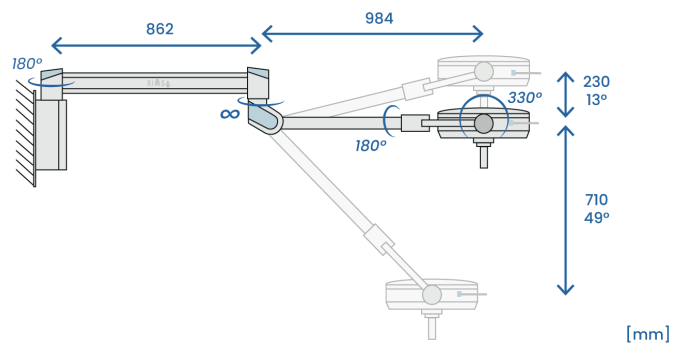


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Wall-mounted version

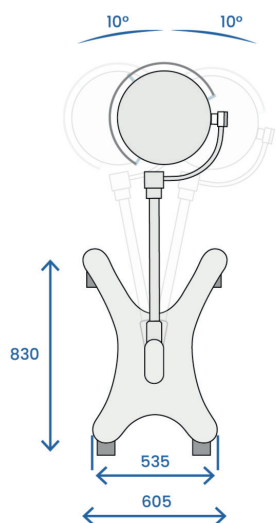


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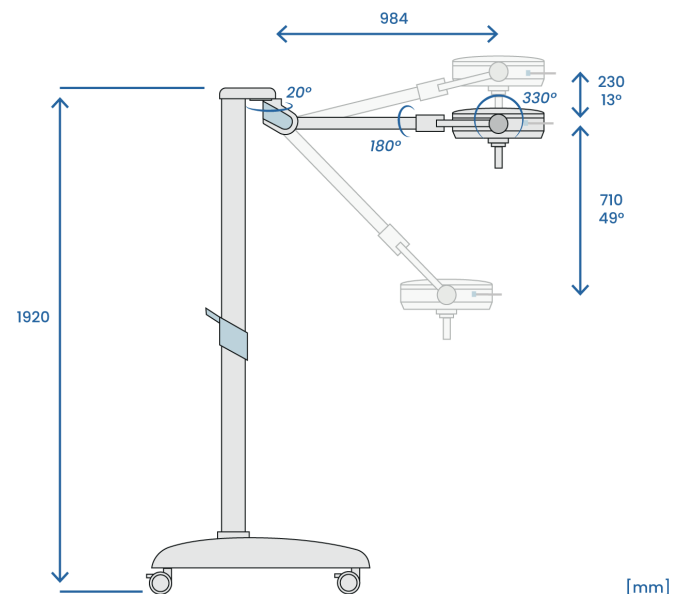


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Mobile version



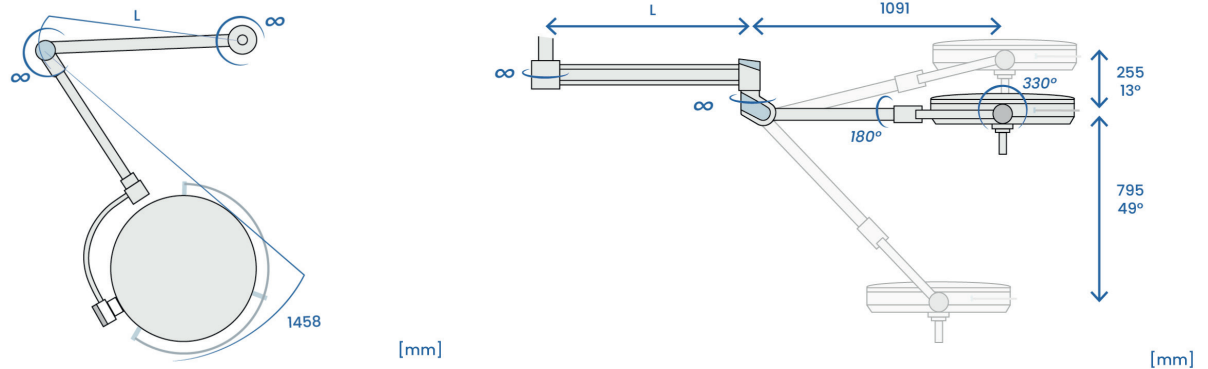
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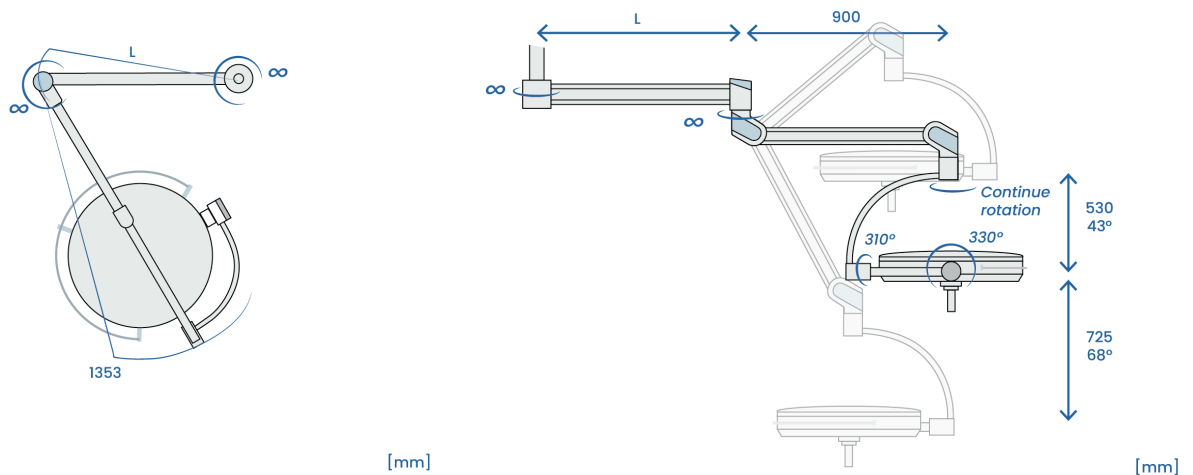
[mm]

Movement PENTALED 63N

Ceiling version with single yoke



Ceiling version with double yoke



6.5 Wall-mounted and mobile version: storage for non-use

If the Product is not used for a longer period of time, set the green switch to the off position (O) and disconnect the power plug from the socket, coil the cable and position it so that it does not get in the way. The environment must not be dusty, damp or exposed to the sun.

Wall-mounted version

Position the horizontal and swinging arms so as not to be in the way of personnel, if possible close to walls.

Mobile version

Move the swinging arm downwards, place the Product in an area that is as free of passage as possible and on a non-tilted surface, and lock the wheels.

7 Product with battery

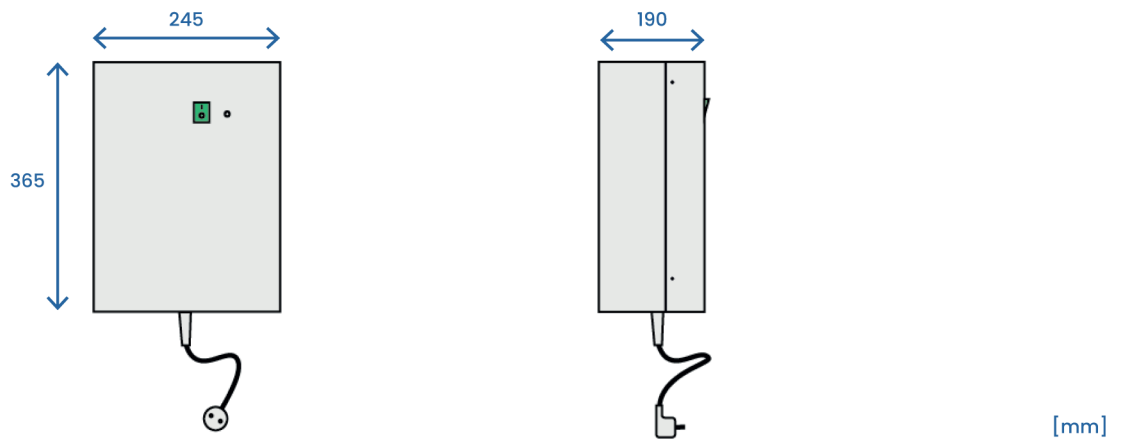
7.1 Description

Upon request, it is possible to equip each lighthouse with a battery pack to allow operation even in the event of a power failure on the mains line.

In the case of a ceiling or wall-mounted Product with battery, the electrical panel and battery pack are supplied together in an additional box (one box for each Product lighthouse).

The system allows the Product to operate either from the mains (100–240 V) or, if the latter is not available, from the 24 V battery. Switching from mains to battery takes place automatically when an electrical fault occurs. To reactivate operation with the mains, the mains must be reactivated. The Product will then automatically reactivate operation with the mains.

Dimensions



Orientation

The electrical panel with battery can be positioned in any orientation. Take care that the I/O switch is accessible.

Power

- INPUT: 100–240 Vac
- OUTPUT: 24 Vdc

Specifications

- Nominal voltage: 12 V
- Capacity: 12 Ah
- Maximum charging current: 3 A
- Weight: 3.75 kg
- Dimensions: 151x98x94 mm
- Operating range: -20 °C / +40 °C

Storage Warnings

- The battery packs are supplied with two charged batteries connected in series with each other.
- Store the batteries at a temperature between -20 °C and +40 °C.
- During storage, recharge the batteries at least once every 6 months.
- The battery ages during storage, so it is recommended to use it as soon as possible.

7.2 Battery life

The estimated battery life is as follows:

Lamp model	Duration [h]
PENTALED 30E	4
PENTALED 30N	4
PENTALED 63N	1.5

7.3 Charging cycle

Connect the battery pack to the mains to charge the batteries. At least 4 hours are required for a full charge. Battery charge is indicated by a bright coloured LED:

- if the light is red, it means that the battery is charging;
- if the light is yellow, it means that the battery is charged between 80% and 95%;
- if the light is green, the battery is fully charged.

When the battery pack is charging, the LED is lit.

If the power supply is interrupted, the LED goes out.

The number of battery life cycles depends on the charge/discharge cycle used:

- if the batteries are fully discharged, the number of theoretical life cycles is 250;
- if the batteries are discharged by 50% and subsequently recharged, the number of theoretical life cycles is 550;
- if the batteries are discharged by 30% and subsequently recharged, the number of theoretical life cycles is 1200.

7.4 Recommendations

It is recommended to perform a battery charge/discharge cycle at least once a month:

- charge the batteries for at least 6 hours
- leave the device switched on until the batteries are discharged;
- if the batteries do not meet the estimated service life, it is recommended that they be replaced.



To ensure optimal operation of the batteries, they should be replaced at least every 3 years.

7.5 Warnings for use

It is recommended to carry out monthly test runs to check the function of the changeover relay.

To do this, disconnect the Product from the power supply and check that the switchover to battery power takes place automatically (the Product must not switch off).



If the Product is not in use and is disconnected from the power supply, set the switch to the off (O) position to prevent the batteries from being discharged.



If the Product is not used and is disconnected from the power supply for more than a month, set the switch to the off (O) position and disconnect the electrical bridge between the two batteries.

7.6 Maintenance

The battery pack does not require any ordinary maintenance. However, once a year, it is necessary to clean its interior with compressed air to remove dust from the various components, thus ensuring correct operation. On this occasion, it is also necessary to check the tightness of the terminals.

8 Cleaning and disinfection

The RESPONSIBLE ORGANISATION must comply with the prescriptions (standards and guidelines) on hygiene, disinfection and sterilisation laid down by the competent national commission.

8.1 Application method



Before cleaning/disinfecting the Product, disconnect the power supply.



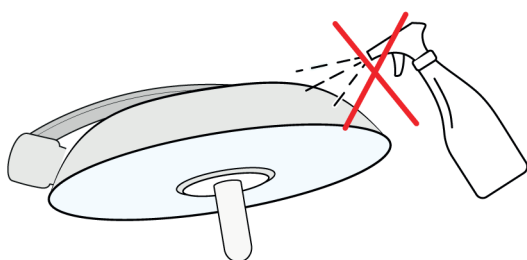
Allow the Product to cool down and only proceed with cleaning/disinfection when it is completely cold.



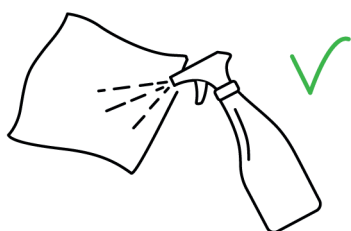
Possible damage to the Product.

To avoid damaging the Product, follow the instructions in this section.

These instructions apply to the cleaning/disinfection of all Product parts.

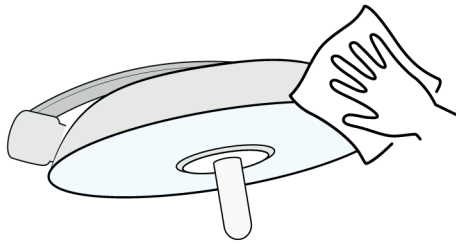


Do not spray detergent/disinfectant or other liquids directly onto the Product.



Dampen a cloth with the cleaner/disinfectant without saturating it.

It is recommended not to use a paper cloth as, once dampened, it may crumble and some fragments may enter the device's crevices.



Wipe the cloth over the Product.

Non-compliance with the requirements described above could result in:

- the detachment of paint with the possibility of it accidentally falling into the operating field;
- the premature ageing of plastics with consequent weakening and possibility of breakage;
- the dulling of protective shields and glass panes.

8.2 Product Cleaning

Frequency

We recommend cleaning the Product daily.



Possible damage to the Product.

To avoid damaging the Product, follow the instructions in this section.

- Do not use sharp, pointed or abrasive objects to avoid the risk of damaging surfaces.
- Do not spill liquids directly onto the Product.
- Clean the Product with a damp but not wet cloth.
- Use appropriate low-alkaline and chlorine-free cleaning agents. Do not use abrasive and/or harmful products (e.g. petrol, paint thinners, alkaline, acidic or alcohol-containing cleaners or aldehydes).
- Dose the cleaning agents scrupulously observing the percentage indications in the manufacturer's technical data sheet, taking care that no liquids penetrate into the crevices of the various parts of the Product, paying particular attention to the lighthouse and support structure.

8.3 Product Disinfection

Frequency

We recommend disinfecting the Product before each use.

Disinfectants may contain substances that are harmful to health; use disinfectants established by the competent national commission for hygiene and disinfection, in compliance with the hygiene standards adopted by the RESPONSIBLE ORGANISATION.



Possible damage to the Product.

To avoid damaging the Product, follow the instructions in this section.

- Do not use sharp, pointed or abrasive objects to avoid the risk of damaging surfaces.
- Do not spill disinfectant liquids directly onto the Product.
- Disinfect the Product with a damp but not wet cloth.
- Use appropriate low-alcohol disinfectants.
- To avoid damage to stainless steel and aluminium parts, use only disinfectants that do not contain chlorine or halogens.

- Dilute the disinfectants scrupulously observing the percentage indications given in the manufacturer's technical data sheet, taking care that no liquid penetrates into the cracks of the various parts of the Product, paying particular attention to the lighthead and support structure.

8.4 Cleaning, disinfection and sterilisation of the handle

Frequency

The handle must be sterilised before use and can withstand about 200 sterilisation cycles.



Danger to the patient.

The OPERATOR must comply with the requirements laid down by the competent national commission for hygiene, disinfection and sterilisation.

The handle is made of polysulphone (PSU), a heat- and impact-resistant plastic material. Replace the handle as soon as it is cracked or deformed, as it could fall into the operating field.

Mounting/dismounting the handle

- Assembly: Slide the handle onto the holder and rotate it until the steel lever clicks into its original position and the rotation is locked. Finally, make sure that the handle is firmly secured.
- Disassembly: press the handle release lever and pull it out.

Cleaning

Before disinfection and sterilisation, clean the handle. It is recommended to use medium alkaline detergents free of active chlorine. Dose the detergents strictly according to the percentage specifications on the manufacturer's data sheet.

Both manual and automated cleaning are permitted.

- For manual cleaning, soak the handle in cleaning solution and clean it with a soft brush or lint-free cloth. After cleaning, rinse off the detergent residue with plenty of water.
- For automated cleaning, use a machine/scrubber that complies with the requirements of ISO 15883-1.

Disinfection

Before sterilisation, disinfect the handle. It is recommended to use alcohol- or aldehyde-based products. Disinfectants must be approved by the disinfectant manufacturer for use on Polysulphone (PSU). Dilute disinfectants strictly according to the percentage specifications on the manufacturer's data sheet.

Both manual and automated disinfection are allowed.

- For manual disinfection, follow the instructions on the manufacturer's data sheet/instructions for use. After disinfection, rinse disinfectant residues thoroughly with water.
- For automated disinfection, use a washer-disinfector complying with the requirements of ISO 15883-1.

Sterilisation

Polysulphone (PSU) is tested for gamma-ray, steam and ethylene oxide (ETO) sterilisation procedures. The result of other sterilisation procedures is not guaranteed.

RIMSA, based on the design of the handle and its own experience, suggests steam sterilisation.

Before sterilising the handle, place it in a suitable sterilisation pack (e.g. plastic/paper bags; single or double pack).

When inserted into the autoclave, take care that the open side of the handle is facing downwards.

The handle must be free and must not be encumbered by other material to be sterilised.

The handle can withstand about 200 steam sterilisation cycles while respecting the following parameters:

- steam sterilisation at 121 °C and 1.3 bar for 25 to 30 minutes;
- steam sterilisation at 134 °C and 2.3 bar for 4 minutes.



Danger to the patient.

Do not exceed the sterilisation temperature of 134 °C. Damaged handles must not be used. Strictly follow ISO 17665-1 and ISO 17665-2.

9 Adjustment and maintenance

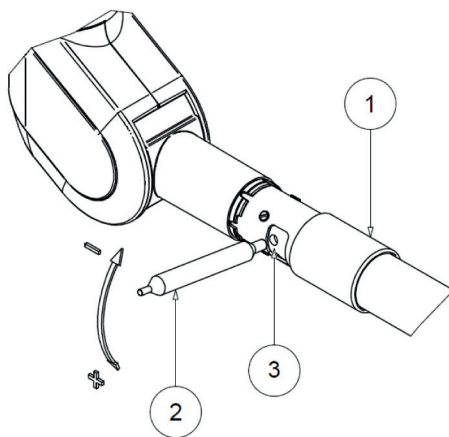
9.1 Swinging/pantograph arm adjustment

If necessary, the pantograph arm can be adjusted:

- with respect to its balance position;
- with respect to its degree of upward swinging.

Adjusting the balance position

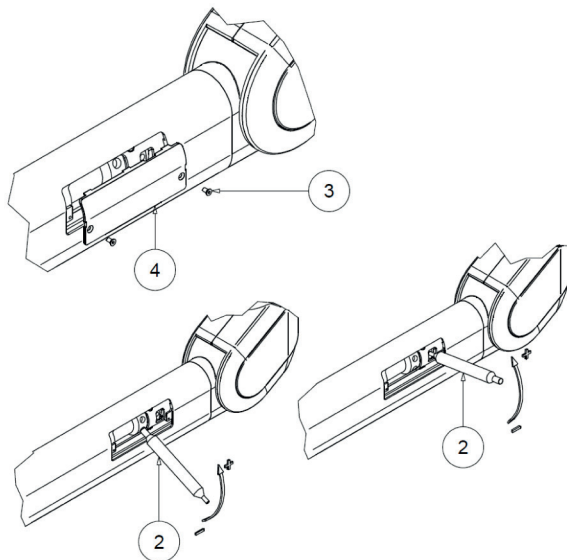
The Product is sold already balanced and requires no further adjustment. Should the spring-balanced swinging/pantograph arm stiffen or loosen over time, it is possible to intervene mechanically by adjusting the compression of the internal spring.



In the case of swinging arms (for single yoke): Manually move the cover (1) forwards. Looking at the lamp from the front, insert the adjustment pin (2) into the holes of the ring nut (3) on the left side of the arm.

- If the swinging arm tends to sag, the spring force is insufficient: turn the ring nut downwards to load the spring.
- If the swinging arm tends to swing upwards, the spring force is too high: turn the ring nut upwards to relieve the spring.

At the end of the adjustment manually reposition the cover (1) to its original position.



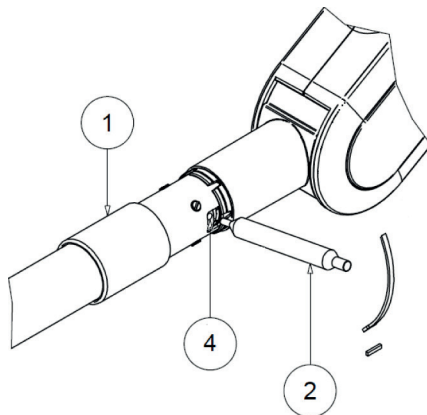
In the case of a pantograph arm (for double yoke): To access the adjustment ring nut, remove screws (3) and cover (4). Insert the adjustment pin (2) into the holes of the coarse adjustment ring nut.

- If the swinging arm tends to sag, the spring force is insufficient: turn the ring nut upwards to load the spring.
- If the swinging arm tends to swing upwards, the spring force is too high: turn the ring nut downwards to relieve the spring.

When the adjustment is complete, manually reposition the cover (4) to its original position and fasten it with the screws (3).

Upward swinging adjustment

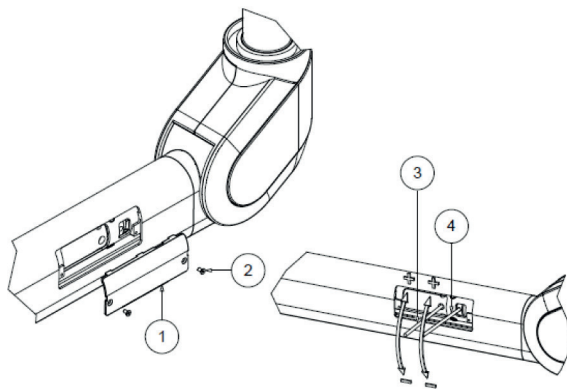
Depending on the size of the installation room (e.g. particularly low ceilings), it may be necessary to limit or increase the upward swinging/pantograph arm's heeling. The downward swinging cannot be modified.



In the case of swinging arms (for single yoke): Manually move the cover (1) forwards. Looking at the lamp from the front, insert the adjusting pin (2) into the holes of the ring nut (4) on the right side of the arm.

- If it is necessary to decrease the upward swinging, turn the ring nut downwards.
- If it is necessary to increase the upward swinging, turn the ring nut upwards.

At the end of the adjustment manually reposition the cover (1) to its original position.

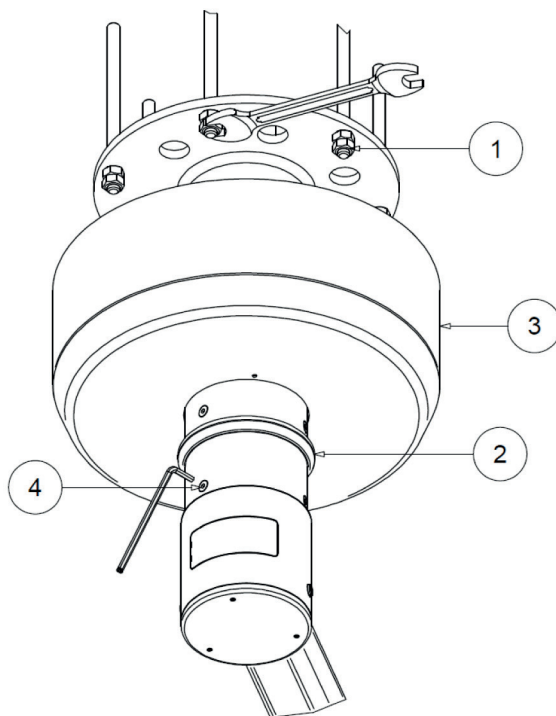


In the case of a pantograph arm (for double yoke): To access the adjustment ring nut, remove screws (3) and cover (4). Insert the adjustment pin (2) into the holes of the smaller adjustment ring nut.

- If it is necessary to decrease the upward swinging, turn the adjustment ring nut downwards.
- If it is necessary to increase the upward swinging, turn the ring nut upwards.

When the adjustment is complete, manually reposition the cover (4) to its original position and fasten it with the screws (3).

9.2 Clutch adjustment



The clutches are adjusted during installation. As with all mechanical parts, the clutches are also subject to wear. If the Product does not remain in a stable position, the clutches involved must be adjusted.

Use a slotted screwdriver to increase the braking force by turning clockwise:

- the screws (1) on the pin of the horizontal arm structure;
- the screws (2) on the swinging/pantograph arm pivot;
- the screws (3) on the single yoke attachment or the screws (5) and (6) on the double yoke attachment;
- the screws (4) or (7) on the lighthouse attachment.

9.3 Checks to be carried out on Product before use



Carry out the electrical verification of the Product.

Before putting into service, carry out the electrical tests and the requirements specified in IEC 62353.

Before each use, in order to ensure Product safety and proper functionality, the OPERATOR must:

- clean/disinfect the Product in accordance with the provisions laid down by the competent national commission;
- check that the light emitted is stable and of adequate intensity;
- check that the lighthouse holds its position correctly;
- check that the swinging/pantograph arm holds its position correctly;
- check that there are no pieces or fragments of paint that can detach and fall onto the operating field. If they are present, remove them manually;
- check that the light source's protective shields are not damaged.

If one or more of these requirements are not met, please contact customer service.

9.4 Ordinary maintenance



Disconnect the power supply before carrying out any maintenance work,



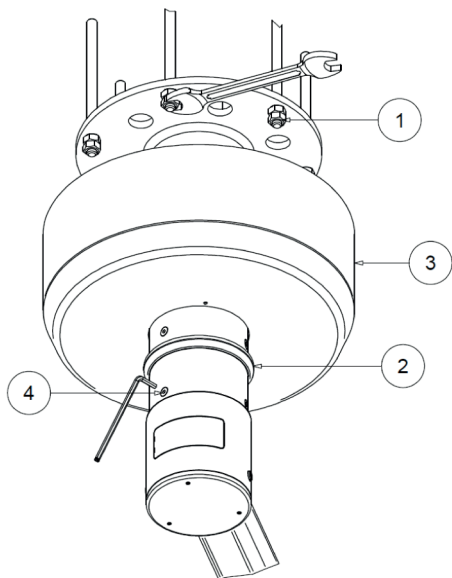
Check the integrity of the Product.



No modification of the Product is permitted.

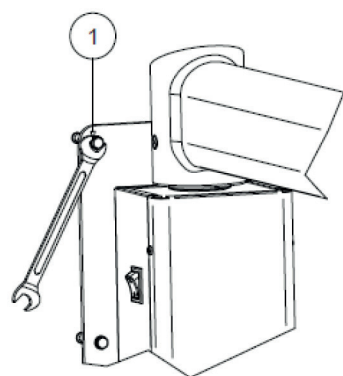
Once a year	Carry out a complete revolution of all the Product's joints and check that there are no noises or squeaks. If this is not the case, lubricate the clutches concerned with industrial grease suitable for use at a service temperature of -30 °C to +120 °C, type OKS 470 or similar.
	If the Product does not maintain its balanced position, adjust the swinging/pantograph arm as indicated in the relevant section.
	If the Product does not maintain a stable position, adjust the clutches as indicated in the relevant section.

Once a year (ceiling version only)



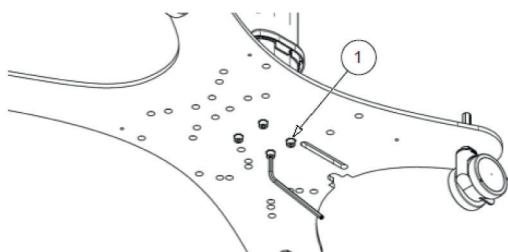
Check that the tiges' fastening nuts (1) are securely tightened. To access the nuts (1), loosen the tiges cover retaining ring screws, then lower the tiges cover. Also check the anchor screws (4) of the horizontal arm to the tiges. If they are loose, tighten them carefully.

Once a year (wall-mounted version only)



Check that the wall mounting screws (1) are properly tightened. If they are loose, tighten them properly.

Once a year (mobile version only)



Check that the stem fixing screws (1) below the crankcase are properly tightened. If they are loose, tighten them appropriately.

9.5 Troubleshooting



Possible damage to the Product.



Presence of dangerous voltage.



No modification of the Product is permitted.

N°	Problem	Solution
1	Product does not remain in a stable position	Check whether the instructions in the installation manual have been followed. Refer to the adjustment instructions in the user and maintenance manual.
2	Light flickers or produces a stroboscopic effect	Contact customer service.
3	Product does not switch on or does not function properly	Check the supply voltage, check the presence and condition of the fuses in the electrical panel, check the connection of the electrical connectors outside and inside the Product, check the voltage continuity inside the Product. If the problem persists, contact customer service.
4	The fuse continues to burn	Check the characteristics of the inserted fuses.

Fuse Replacement

The electrical panel must be accessed for fuse replacement. Once you have identified the location of the electrical panel, follow the instructions below.

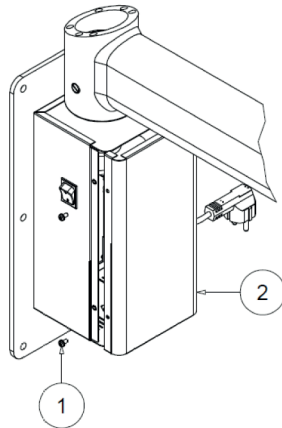


Disconnect voltage before performing any operation.

In the case of electrical panels mounted on tiges:

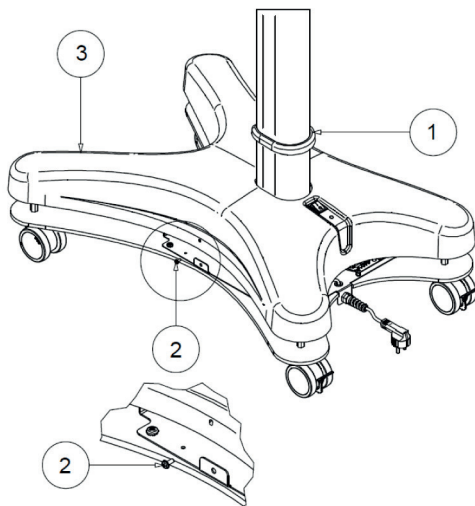
Loosen the screws of the tiges cover, then lower the tiges cover so that the electrical panel can be accessed.

Remove the screw fuse holder from the terminal box and replace the fuse, taking care to replace it with one of the same type.



In the case of wall-mounted version: remove the 4 screws (1) and remove the cover (2) of the box with electrical panel.

Remove the screw fuse holder from the terminal box and replace the fuse, taking care to replace it with one of the same type.



In the case of the mobile version: Lift up the ring (1), remove the screws (2) and lift off the cover (3).

Remove the screw fuse holder from the terminal box and replace the fuse, taking care to replace it with one of the same type.

If necessary, RIMSA shall provide wiring diagrams, parts lists, descriptions, adjustment instructions, or other information to assist the TECHNICAL SERVICE PERSONNEL in repairing those parts of the Product indicated as repairable by the TECHNICAL SERVICE PERSONNEL.

If the above indications are not sufficient to solve the problem, please contact customer service.

9.6 Repair and extraordinary maintenance



The Product must only be opened and repaired by TECHNICAL SERVICE PERSONNEL.



Switch off the power supply before carrying out any repair/maintenance work.

In the event of Product malfunction or failure, contact the manufacturer's customer service department. If necessary, the manufacturer will provide wiring diagrams, parts lists, descriptions, operating instructions or other information to assist the TECHNICAL SERVICE PERSONNEL in repairing the Product.



Carry out the electrical verification of the Product.

After every repair and/or extraordinary maintenance, carry out electrical tests and the requirements specified in IEC 62353.



No modification of the Product is permitted.

9.7 Spare parts list



Use only original spare parts.

Description	Order code
Fuse T2AH 250V '5x20'	Z400195
Fuse T4AH 250V '5x20'	Z400215
Fuse T10AH 250V '5x20'	Z400217
Sterilizable handle	Z200518
Switching power supply SERIES E/N	Z400629
Electronic board SERIES E/N	Z300632
I/O switch (for wall-mounted and mobile versions)	Z300016
Membrane keyboard SERIES E	Z300227
Membrane keyboard SERIES N	Z300234

9.8 Disposal at end of use

Comply with current waste disposal regulations. This Product must not be disposed of in normal waste bins. To avoid risks to the environment and health resulting from the dispersion of pollutants into the environment, separate the various internal components (e.g. iron, aluminium, plastic and electrical material) and take them to the appropriate centres for proper recycling.

10 Technical data

10.1 Technical data PENTALED 30E

Photometric data	
Color temperature, CCT $\pm 10\%$ [K]	4500 / 5000
Illuminance at center $E_{C,Ref} \pm 10\%$ [lux]	160 000
Reference distance D_{Ref} [cm]	100
Max illuminance at center $E_{C,Mi} \pm 10\%$ [lux]	160 000
Max illumination distance D_{Mi} [cm]	100
Light field diameter $d_{10} \pm 10\%$ [mm]	220
Light field diameter $d_{50} \pm 10\%$ [mm]	130
$d_{50} / d_{10} [\%]$	≥ 50
Depth of illumination $L1+L2$ at 60% $\pm 10\%$ [mm]	645
Remaining illuminance $E_{C,Ref}$ with 1 mask $\pm 5 [\%]$	0
Remaining illuminance $E_{C,Ref}$ with 2 masks $\pm 5 [\%]$	42
Remaining illuminance $E_{C,Ref}$ with cavity $\pm 5 [\%]$	100
Remaining illuminance $E_{C,Ref}$ with cavity and 1 mask $\pm 5 [\%]$	0
Remaining illuminance $E_{C,Ref}$ with cavity and 2 masks $\pm 5 [\%]$	42
Color rendering index $R_a \pm 5 [-]$	95
$R_g \pm 5 [-]$	90
Total irradiance $E_{Total} (D_{Ref}) \pm 10\% [W/m^2]$	627
Total irradiance $E_{Total} (D_{Mi}) \pm 10\% [W/m^2]$	627
Electrical connection data	
Primary alternating voltage [Vac]	100 – 240
Frequency [Hz]	50/60
Power consumption [VA]	60
Power consumption with fully discharged battery [VA]	150

Light source	n. 30 LED
LED light source lifespan [h] (this data may vary based on voltage spikes and usage frequency)	60 000

General data	
Regulation	REGULATION (EU) 2017/745
Standards	IEC 60601-1, IEC 60601-1-2, IEC 60601-2-41
Product classification Medical Device	Class I
Operating conditions	Continuous operation
Means of electrical isolation from line voltage	Main switch external to the product for ceiling versions. Main switch on the Product for mobile and wall-mounted versions.
Steam sterilization of the handle	121 °C 1.3 bar from 25 to 30 minutes. 134 °C 2.3 bar for 4 minutes.
Dimensions	
Lamp body diameter [cm]	40
Light emission surface area [cm ²]	762
Product weight single ceiling - double ceiling - wall-mounted - mobile version - mobile version with battery [kg]	39 - 63 - 27 - 47 - 57
Markings	
CE	Compliant with REGULATION (EU) 2017/745
<p><i>Note 1: unless otherwise indicated, all values are to be considered with a tolerance of $\pm 5\%$ due to metrological and construction reasons.</i></p> <p><i>Note 2: all tests have been performed in accordance with section 201.5.4 "General requirements for tests on EM EQUIPMENT - Other conditions" of the IEC 60601-2-41 standard. In particular, the Product was tested with maximum light intensity, color temperature 5000K, and maximum light diameter.</i></p> <p><i>Note 3: the optical radiation emitted by this Product complies with the exposure limits for reducing photobiological risk as per the IEC 60601-2-41 standard.</i></p>	

10.2 Technical data PENTALED 30N

Photometric data	
Color temperature, CCT $\pm 10\%$ [K]	4500 / 5000
Illuminance at center $E_{C,Ref} \pm 10\%$ [lux]	160 000
Reference distance D_{Ref} [cm]	100
Max illuminance at center $E_{C,Ml} \pm 10\%$ [lux]	160 000
Max illumination distance D_{Ml} [cm]	100
Light field diameter $d_{10} \pm 10\%$ [mm]	205
Light field diameter $d_{50} \pm 10\%$ [mm]	125
$d_{50} / d_{10} [\%]$	≥ 50
Depth of illumination L1+L2 at 60% $\pm 10\%$ [mm]	700
Remaining illuminance $E_{C,Ref}$ with 1 mask $\pm 5 [\%]$	0
Remaining illuminance $E_{C,Ref}$ with 2 masks $\pm 5 [\%]$	45
Remaining illuminance $E_{C,Ref}$ with cavity $\pm 5 [\%]$	100
Remaining illuminance $E_{C,Ref}$ with cavity and 1 mask $\pm 5 [\%]$	0
Remaining illuminance $E_{C,Ref}$ with cavity and 2 masks $\pm 5 [\%]$	45
Color rendering index $R_a \pm 5 [-]$	95
$R_g \pm 5 [-]$	90
Total irradiance $E_{Total} (D_{Ref}) \pm 10\% [W/m^2]$	580
Total irradiance $E_{Total} (D_{Ml}) \pm 10\% [W/m^2]$	580
Electrical connection data	
Primary alternating voltage [Vac]	100 – 240
Frequency [Hz]	50/60
Power consumption [VA]	60
Power consumption with fully discharged battery [VA]	150
Light source	n. 30 LED

LED light source lifespan [h] (this data may vary based on the occurrence of voltage spikes and frequency of use)	60 000
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General Data	
Regulation	REGULATION (EU) 2017/745
Standards	IEC 60601-1, IEC 60601-1-2, IEC 60601-2-41
Product classification Medical Device	Class I
Operating conditions	Continuous operation
Means of electrical isolation from line voltage	Main switch external to the Product for ceiling versions. Main switch on the Product for mobile and wall-mounted versions.
Steam sterilization of the handle	121 °C 1.3 bar from 25 to 30 minutes. 134 °C 2.3 bar for 4 minutes.
Dimensions	
Lamp body diameter [cm]	40
Light emission surface [cm ²]	762
Product weight single ceiling - double ceiling - wall - mobile - mobile battery [kg]	39 - 63 - 27 - 47 - 57
Markings	
CE	Compliant with REGULATION (EU) 2017/745
<p><i>Note 1: unless otherwise indicated all values are to be considered with a tolerance of $\pm 5\%$ due to metrological and construction reasons.</i></p> <p><i>Note 2: all tests were conducted in accordance with section 201.5.4 "General requirements for tests on EM EQUIPMENT - Other conditions" of the IEC 60601-2-41 standard. Specifically, the Product was tested with maximum luminous intensity, color temperature 5000K, and minimum luminous diameter.</i></p> <p><i>Note 3: the optical radiation emitted by this Product complies with the exposure limits for reducing photobiological risk as per the IEC 60601-2-41 standard.</i></p>	

10.3 Technical data PENTALED 63N

Photometric Data	
Color temperature, CCT $\pm 10\%$ [K]	4500 / 5000
Illuminance at center $E_{C,Ref} \pm 10\%$ [lux]	160 000
Reference distance D_{Ref} [cm]	100
Max illuminance at center $E_{C,Ml} \pm 10\%$ [lux]	160 000
Max illuminance distance D_{Ml} [cm]	100
Light field diameter $d_{10} \pm 10\%$ [mm]	210
Light field diameter $d_{50} \pm 10\%$ [mm]	110
$d_{50} / d_{10} [\%]$	≥ 50
Depth of illumination L1+L2 at 60% $\pm 10\%$ [mm]	475
Remaining illuminance $E_{C,Ref}$ with 1 mask $\pm 5 [\%]$	80
Remaining illuminance $E_{C,Ref}$ with 2 masks $\pm 5 [\%]$	48
Remaining illuminance $E_{C,Ref}$ with cavity $\pm 5 [\%]$	100
Remaining illuminance $E_{C,Ref}$ with cavity and 1 mask $\pm 5 [\%]$	80
Remaining illuminance $E_{C,Ref}$ with cavity and 2 masks $\pm 5 [\%]$	48
Color rendering index $R_a \pm 5 [-]$	95
$R_g \pm 5 [-]$	90
Total irradiance $E_{Total} (D_{Ref}) \pm 10\% [W/m^2]$	590
Total irradiance $E_{Total} (D_{Ml}) \pm 10\% [W/m^2]$	590
Electrical connection data	
Primary alternating voltage [Vac]	100 – 240
Frequency [Hz]	50/60
Power consumption [VA]	70
Power consumption with fully discharged battery [VA]	160
Light source	n. 72 LED

LED light source duration [h] (this data may vary based on the occurrence of voltage spikes and frequency of use)	60 000
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General data	
Regulation	REGULATION (EU) 2017/745
Standards	IEC 60601-1, IEC 60601-1-2, IEC 60601-2-41
Product classification Medical Device	Class I
Operating conditions	Continuous operation
Means of electrical isolation from line voltage	Main switch external to the Product for ceiling versions. Main switch on the Product for mobile and wall-mounted versions.
Steam sterilization of the handle	121 °C 1.3 bar from 25 to 30 minutes. 134 °C 2.3 bar for 4 minutes.
Dimensions	
Diameter of lamp body [cm]	63
Light emission surface [cm ²]	1828
Product weight single ceiling - double ceiling - wall - mobile - mobile battery [kg]	47 - 79 - / - / - /
Markings	
CE	Compliant with REGULATION (EU) 2017/745
<p><i>Note 1: unless otherwise indicated, all values are to be considered with a tolerance of $\pm 5\%$ due to metrological and construction reasons.</i></p> <p><i>Note 2: all tests were conducted in accordance with section 201.5.4 "General requirements for tests on EM EQUIPMENT - Other conditions" of the IEC 60601-2-41 standard. Specifically, the Product was tested with maximum light intensity, color temperature 4500 K, and minimum luminous diameter.</i></p> <p><i>Note 3: the optical radiation emitted by this Product complies with the exposure limits for reducing photobiological risk as specified by the IEC 60601-2-41 standard.</i></p>	

11 EU Declaration of Conformity

Drafted pursuant to Article 19 and Annex IV of REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC.

Manufacturer: **RIMSA P. LONGONI SRL**

Registered Office Address: **Via Monterosa, 18/20/22 - 20831 - Seregno (MB) - Italia**

Unique Registration Number (SRN): **IT-MF-000009224**

This declaration of conformity is issued under the sole responsibility of the manufacturer.

Basic UDI-DI: **++B880LUMINAIREPM**

Product name and trade name: PENTALED 30E

Model reference:

PENTA30ESO — LAMP MODEL PENTALED 30E CEILING

PENTA30E+30E — LAMP MODEL PENTALED 30E DOUBLE CEILING

PENTA30EPA — LAMP MODEL PENTALED 30E WALL

PENTA30EPI — LAMP MODEL PENTALED 30E MOBILE STAND

Product name and trade name: PENTALED 30N

Product name and trade name: PENTALED 63N

Model reference:

PENTA30NSO — LAMP MODEL PENTALED 30N CEILING

PENTA30N+30N — LAMP MODEL PENTALED 30N DOUBLE CEILING

PENTA30NPA — LAMP MODEL PENTALED 30N WALL

PENTA30NPI — LAMP MODEL PENTALED 30N MOBILE STAND

PENTA63NSO — LAMP MODEL 63N CEILING

PENTA63N+30N — LAMP MODEL PENTALED 63N+30N

PENTA63N+63N — LAMP MODEL PENTALED 63N+63N

Intended use: SCIALITIC LAMP FOR SURGICAL LUMINAIRE

Risk class of the device, according to the regulations of Annex VIII of REGULATION (EU) 2017/745: **CLASS I**

Justification:	Duration:	Short-term (Annex VIII, CHAPTER I, point 1. DURATION OF USE)
	Description:	Non-invasive medical device (Annex VIII, CHAPTER III, point 4. NON INVASIVE DEVICES, paragraph 4.1 Rule 1)
		Active medical device (Annex VIII, CHAPTER III, point 6. ACTIVE DEVICES, paragraph 6.2 Rule 10)

The manufacturer declares that the device complies with REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 on medical devices, amending Directive 2001/83/EC, Regulation (EC) No 178/2002 and Regulation (EC) No 1223/2009 and repealing Council Directives 90/385/EEC and 93/42/EEC and with the following regulations:

IEC 60601-1	(Part 1: General requirements for basic safety and essential performance)
IEC 60601-1-2	(Part 1: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility - Requirements and tests)
IEC 60601-2-41	(Part 2: Particular requirements for the basic safety and essential performance of surgical luminaires for diagnosis)

The conformity assessment procedure of the device is carried out according to premise (60) and Article 52 of REGULATION (EU) 2017/745.

The Quality and Environment System of RIMSA complies with UNI EN ISO 9001, UNI CEI EN ISO 13485, ISO 14001 and ISO/IEC 27001 standards and is certified by IMQ S.p.A. (certificates no. 9120.RMS1, no. 9124.RMS2, no. 0833.2023 and no. 1402.2024).

Name: Paolo Longoni
Position: CEO

RIMSA P. LONGONI S.p.A.
Longoni

12 EMC Declaration

The Product has been tested according to IEC 60601-1-2 to ensure correct electromagnetic compatibility.



Possibility of interference with neighbouring equipment

Portable and mobile communication equipment may affect the Product. The Product should not be used in the vicinity of another device and if use is necessary, the Product must be checked for functionality.

The use of accessories other than those supplied/recommended RIMSA may increase the emission level and lower the immunity level of the device.

The Product is designed for use in the electromagnetic environment described below.

It is the responsibility of the RESPONSIBLE ORGANISATION or OPERATOR to ensure that the Product is used in a compatible environment.

It may happen that the Product, when subjected to irradiation in the range 80 MHz – 1 GHz or to burst radiation, no longer responds to commands, neither for the lamp nor for the camera (if present).

In this case, essential performance will still be guaranteed, but to restore normal operation, it will be necessary to remove power from the main switch.

Immunity tests	Compliance	Electromagnetic environment – directives
RF emissions CISPR 11	Group 1	The Product uses RF energy only for its internal operation. Consequently, its RF emissions are very low and are unlikely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class A	The Product is suitable for use in all rooms except domestic rooms, and those directly connected to a low-voltage public mains supply that supplies buildings used for domestic purposes, provided the following warning is given.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuation/ flicker emissions IEC 61000-3-3	Compliant	Warning: This Product is intended for use by professional medical personnel only. This Product may cause radio interference or may disturb the operation of nearby equipment. It may be necessary to take measures to mitigate such interference, such as reorienting and relocating the Product or shielding the room.
NOTE: The EMISSIONS characteristics of this equipment make it suitable for use in industrial areas and hospitals (CISPR 11 class A). If it is used in a residential ENVIRONMENT (for which CISPR 11 class B is normally required) this equipment might not offer adequate protection to radio-frequency communication services. The user might need to take mitigation measures, such as relocating or re-orienting the equipment.		

Immunity tests	Test level IEC 60601-1-2	Compliance level	Electromagnetic environment - directives
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV at contact ±2, ±4, ±8 kV, ±15 kV in air	± 8 kV at contact ±2, ±4, ±8 kV, ±15 kV in air	It is preferable for the floor to be made of wood, concrete or ceramic tiles. If the floor is covered with synthetic materials, the moisture content must be at least 30%.
Fast-pulsing electrical transients IEC 61000-4-4	± 2 kV for power supply lines	± 2 kV for power supply lines	A mains quality typical of a commercial or domestic environment is recommended.
Shockwaves IEC 61000-4-5	± 0.5 kV, ± 1 kV Differential mode	± 0.5 kV, ± 1 kV Differential mode	A mains quality typical of a commercial or hospital environment is recommended.
Magnetic field at mains frequency (50/60 Hz) IEC 61000-4-8	30 A/m	30 A/m	Magnetic fields at mains frequency must be at levels characteristic of a typical location in a commercial or hospital environment.
Voltage dips, short interruptions and variations on power supply input lines IEC 61000-4-11	10 ms - 0% at 0°, 45°, 90°, 135°, 180°, 225°, 270°, 315° 20 ms - 0% at 0° 500 ms - 70% at 0° 5 s - 0%	10 ms - 0% at 0°, 45°, 90°, 135°, 180°, 225°, 270°, 315° 20 ms - 0% at 0° 500 ms - 70% at 0° 5 s - 0%	A mains power quality typical of a commercial or hospital environment is recommended. If the Product is to be used continuously even in the event of a power failure, connect the lamp to a mains supply capable of providing continuous power or to a battery.
NOTE: U_T is the AC mains voltage before application of the test level.			

Immunity tests	Test level IEC 60601-1-2	Compliance level	Electromagnetic environment - directives
RF conducted IEC 61000-4-6	3 V _{eff} 150 kHz to 80 MHz 6 V ISM frequencies	3 V _{eff} 150 kHz to 80 MHz 6 V ISM frequencies	Portable and mobile RF communication systems must be used no closer than 30 cm (12 inches) to any part of the Product, including cables specified by the manufacturer; observe the recommended separation distance, calculated according to the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = 1.2\sqrt{P}$ from 150 KHz to 80 MHz $d = 1.2\sqrt{P}$ 80 MHz to 800 MHz $d = 2.3\sqrt{P}$ 800 MHz to 2.7 GHz where P is the maximum output power of the transmitter in watts [W], according to the transmitter manufacturer and d is the recommended separation distance in metres [m]. Field strengths of fixed RF transmitters, as determined by an electromagnetic site survey, must be below the compliance level in each frequency range. In the vicinity of equipment marked with the following symbol, interference may occur: 
Radiated RF IEC 61000-4-3	3 V/m 80 MHz to 2.7 GHz	3 V/m 80 MHz to 2.7 GHz	

NOTE 1: At 80 MHz and 800 MHz, 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.

Test frequency [MHz]	Band ^{a)} [MHz]	Service ^{a)}	Modulation ^{b)}	Maximum power [W]	Distance [m]	Immunity test level [V/m]
385	380-390	TETRA 400	Pulse modulation ^{b)} 18 Hz	1,8	0,3	27
450	430-470	GMRS 460, FRS 460	FM ^{c)} ± 5 kHz deviation 1 kHz sine	2	0,3	28
710 745 780	704-787	LTE Band 13, 17	Pulse modulation ^{b)} 217 Hz	0,2	0,3	9
810 870 930	800-960	GSM800/900, TETRA 800, iDEN 820, CDMA 850, LTE Band 5	Pulse modulation ^{b)} 18 Hz	2	0,3	28
1720 1845 1970	1700-1990	GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3, 4, 25; UMTS	Pulse modulation ^{b)} 217 Hz	2	0,3	28
2450	2400-2750	Bluetooth, WLAN, 02.11, b/g/n, RFID 2450, LTE Band 7	Pulse modulation ^{b)} 217 Hz	2	0,3	28
5240 5500 5785	5100-5800	WLAN 802-11 a/n	Pulse modulation ^{b)} 217 Hz	0,2	0,3	9

NOTE: If it is necessary to achieve the IMMUNITY TEST LEVEL, the distance between the transmitting antenna and the ME medical device or ME SYSTEM must be reduced to 1 m. The 1 m test distance is permitted by IEC 61000-4-3.

^{a)} For some services only rising frequencies are included.

^{b)} The carrier must be modulated using a square wave signal at 50% duty cycle.

^{c)} As an alternative to FM modulation, a modulation of 50% of the pulse at 18 Hz may be used because, although it does not represent actual modulation, it would be the worst case.

Recommended separation distance between portable and mobile RF communication equipment and the Product

The Product is intended for use in electromagnetic environments in which radiated RF interference is controlled. The customer or the user of the Product may prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communication equipment (transmitters) and the Product as specified below, according to the maximum output power of the communication equipment.

Maximum emission power assigned to the transmitter [W]	Separation distance as a function of transmitter frequency [m]		
	150 kHz to 80 MHz $d = 1.2\sqrt{P}$	80 MHz to 800 MHz $d = 1.2\sqrt{P}$	800 MHz to 2.7 GHz $d = 2.3\sqrt{P}$
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 with a maximum output power level not listed in the above table, the recommended separation distance in metres [m] can be determined using the equation applicable to the frequency of the transmitter, where P is the maximum output power of the transmitter in watts [W] in accordance with the transmitter manufacturer.

Note 1: At 80 MHz and 800 MHz, the separation distance for the upper frequency range applies.

Note 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflections from the structures of objects and people.

**Proximity immunity to magnetic field in the frequency range 9 kHz - 13.56 MHz
IEC 61000-4-39**

Magnetic fields at mains frequency must be at characteristic levels caused by radio frequency fields from devices used in close proximity.

Test Frequency	Modulation	Immunity test level [A/m]
30 kHz ^{a)}	CW	8
134.2 kHz	Pulse modulation ^{b)} 2.1 kHz	65 ^{c)}
13.56 MHz	Pulse modulation ^{b)} 50 kHz	7.5 ^{c)}

a) This test is only applicable to ME MEDICAL DEVICES or ME SYSTEMS intended for use in a healthcare home environment.

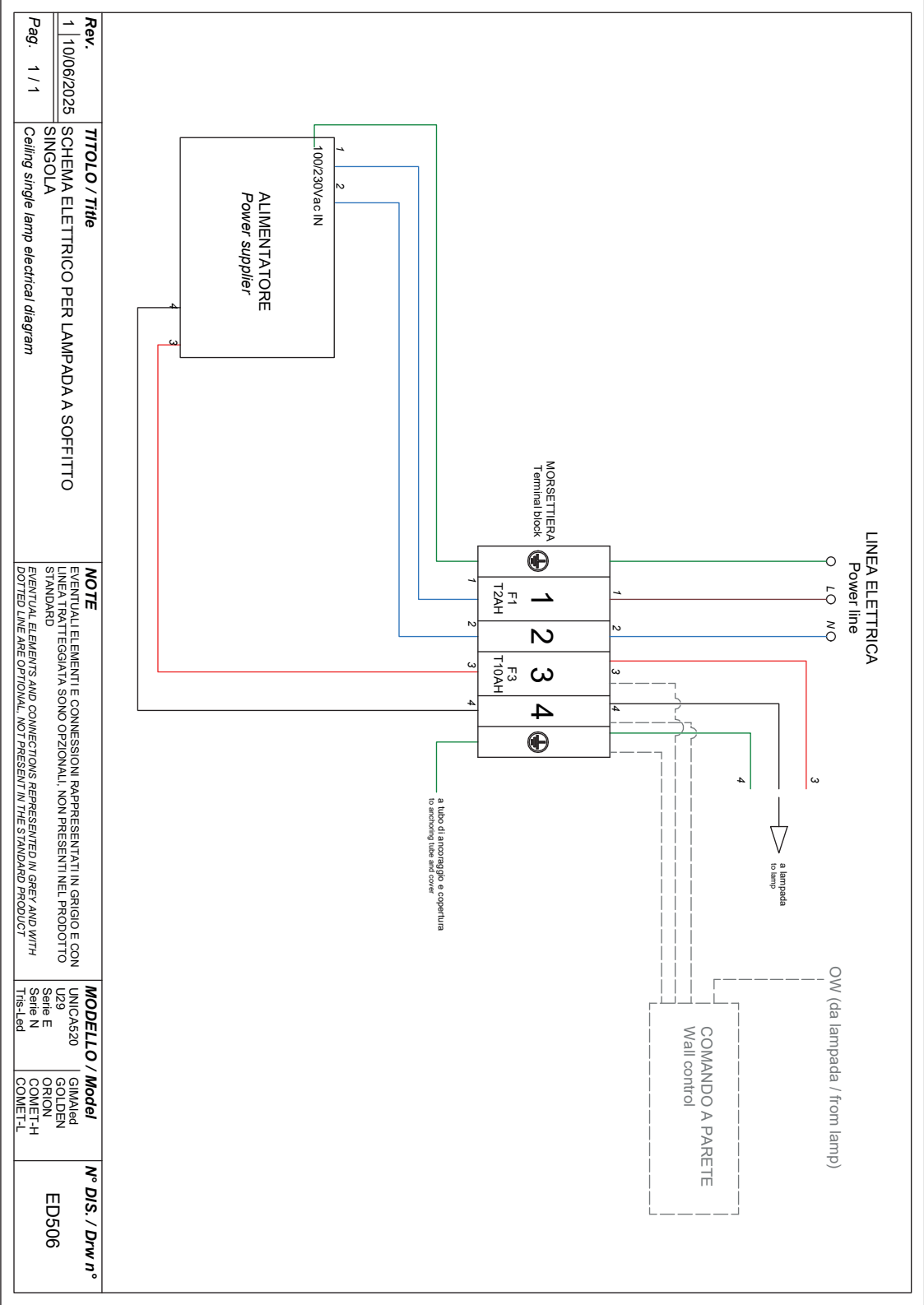
b) The vector shall be modulated using a square wave signal at 50% duty cycle.

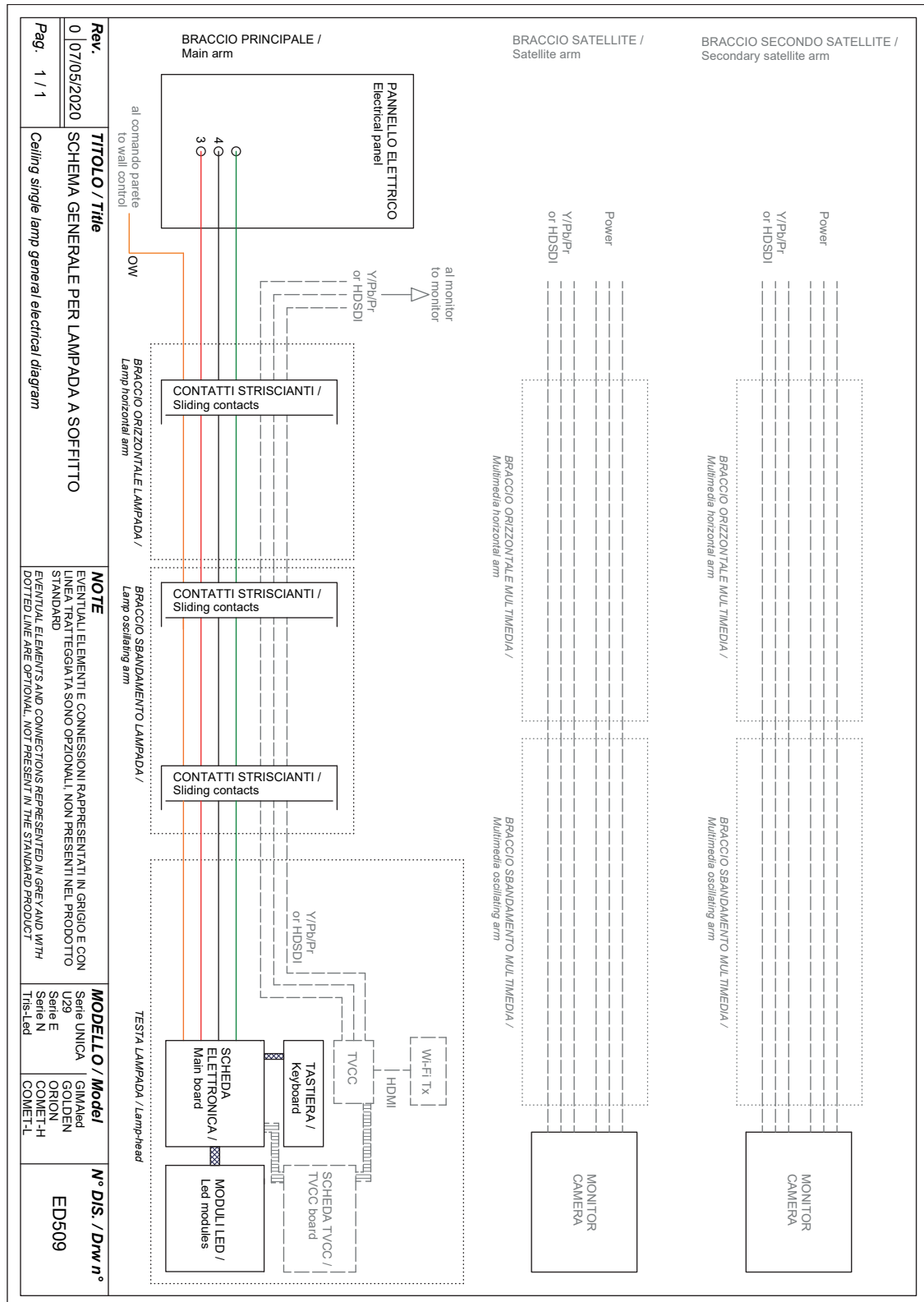
c) The mean square value shall be applied prior to modulation.

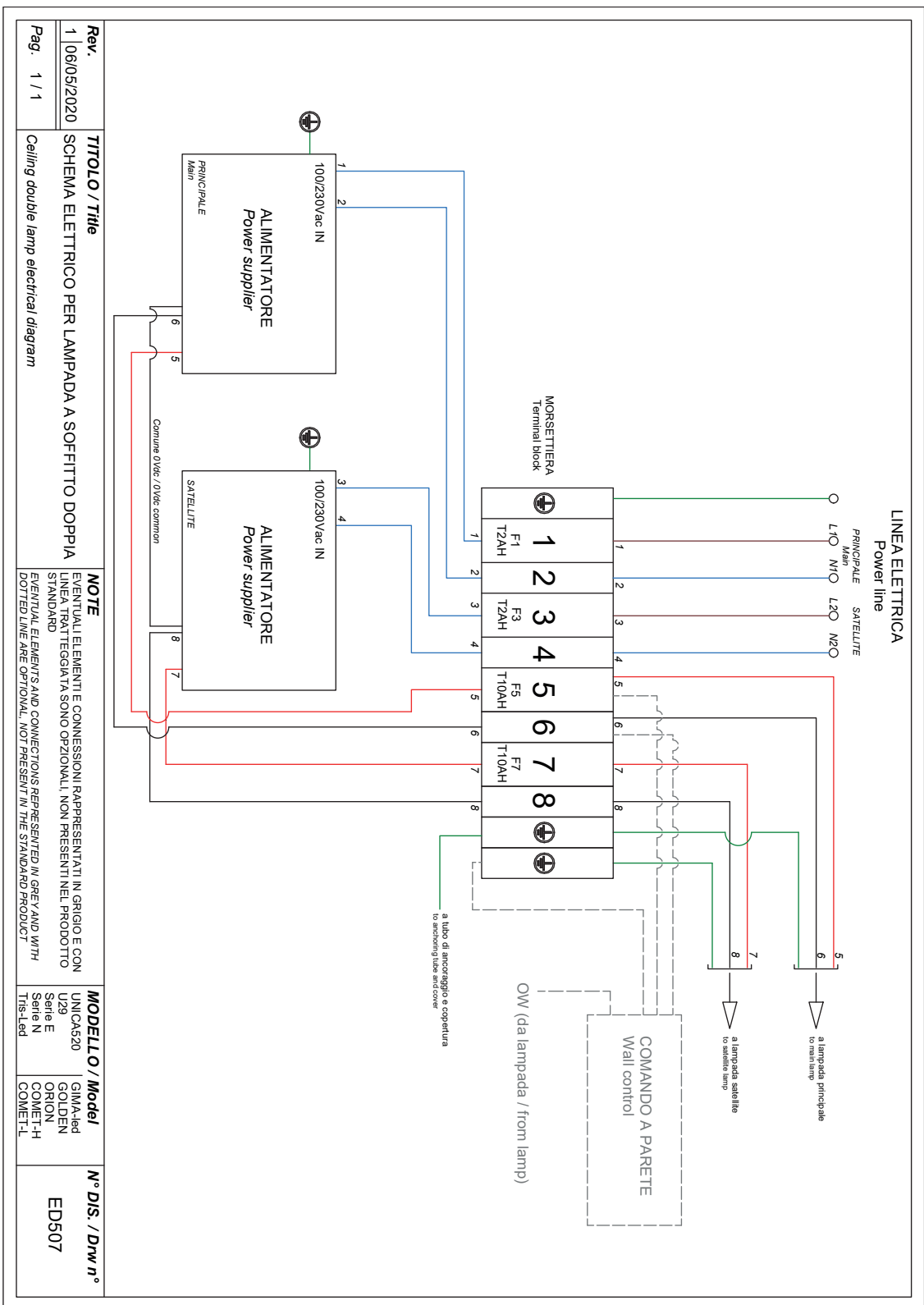
13 Certificate of Warranty

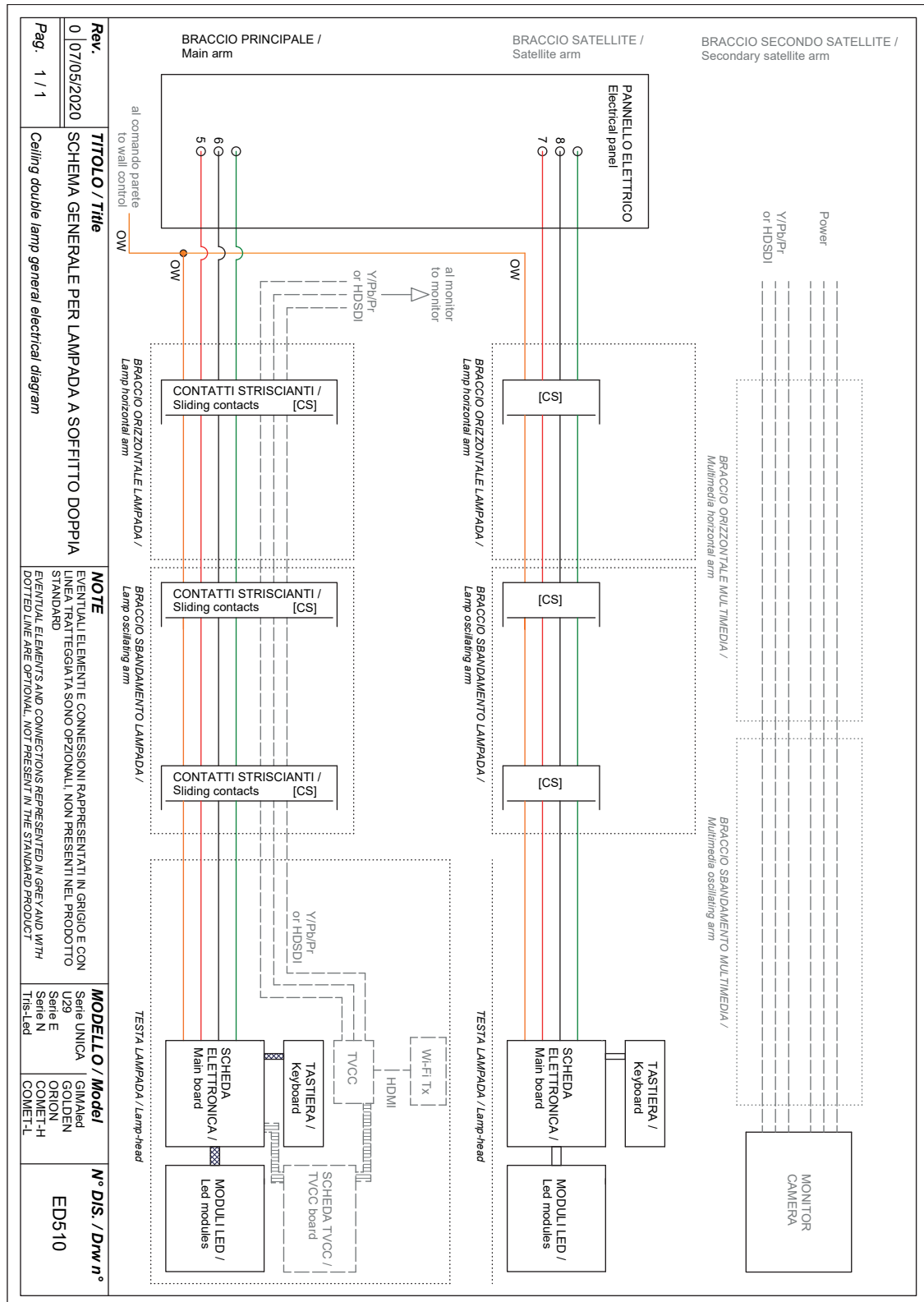
- 1** The Product is under warranty for a period of 18 months, including electrical parts.
- 2** The warranty begins on the date of dispatch of the Product from the RIMSA warehouse to the buyer.
- 3** In the event of a dispute, the date indicated on the 'transport document' accompanying the goods shall be deemed valid.
- 4** The warranty is limited to sending the buyer spare parts of the Product or, if RIMSA deems replacement of the parts to be impracticable, to replacing the entire Product, and is made for well established causes of manufacture and at RIMSA's sole discretion. The warranty does not therefore include any other costs or expenses (such as but not limited to labour costs, packaging and transport costs, etc.).
- 5** Excluded from the guarantee are components subject to normal wear and tear (e.g. halogen bulbs, LEDs, fuses, relays, ball bearings, etc.).
- 6** They are not covered by warranty:
 - malfunctions due to non-compliance with all instructions in the manuals
 - malfunctions due to installation and/or maintenance errors;
 - malfunctions or defects caused by neglect, negligence, misuse or other causes not attributable to RIMSA;
 - malfunctions or defects due to the fact that the electrical installation in the environment (room) in which the installation is carried out does not comply with IEC 60364-7-710 (standard for electrical installations for medical premises) and similar standards.
- 7** RIMSA shall indemnify direct damage to the buyer that is documented as being attributable to his Product, caused within the warranty period, in an amount not exceeding 40% of the net value of the Product as stated on the invoice to the buyer. RIMSA's liability for indirect or consequential damage (including loss of use of the Product) arising from the delivery is expressly excluded.
- 8** The guarantee in this certificate is in lieu of the legal guarantees for defects and non-conformities and excludes any other possible liability of RIMSA arising from the Products supplied.
- 9** Compensation for any damage to persons or property caused by the Product's malfunction or defects shall be limited to the amount of RIMSA's liability insurance coverage.
- 10** The purchaser automatically forfeits the guarantee if:
 - the Product has been tampered with or modified by the buyer or a third party;
 - the Product has been repaired by the Buyer or a third party without complying with the instructions contained in the Manuals
 - the serial number of the Product has been erased, blurred or removed;
 - the buyer is not in good standing with payments.
- 11** For warranty work, the purchaser must only contact RIMSA.
- 12** Parts replaced under warranty must be returned to RIMSA, only if requested by RIMSA, carriage paid and properly packaged.
- 13** Failure to return the component at RIMSA's request will result in the cost of the component being charged.
- 14** RIMSA does not accept returns from end users or otherwise from parties other than the buyer.
- 15** Products returning to RIMSA must have the return authorisation documentation and a document describing the malfunction attached.
- 16** For anything not covered by this warranty certificate, please refer to Italian law.
- 17** For any dispute arising out of or in connection with orders to which this guarantee certificate applies that the parties have not been able to settle amicably, the Court of Milan shall have exclusive jurisdiction.

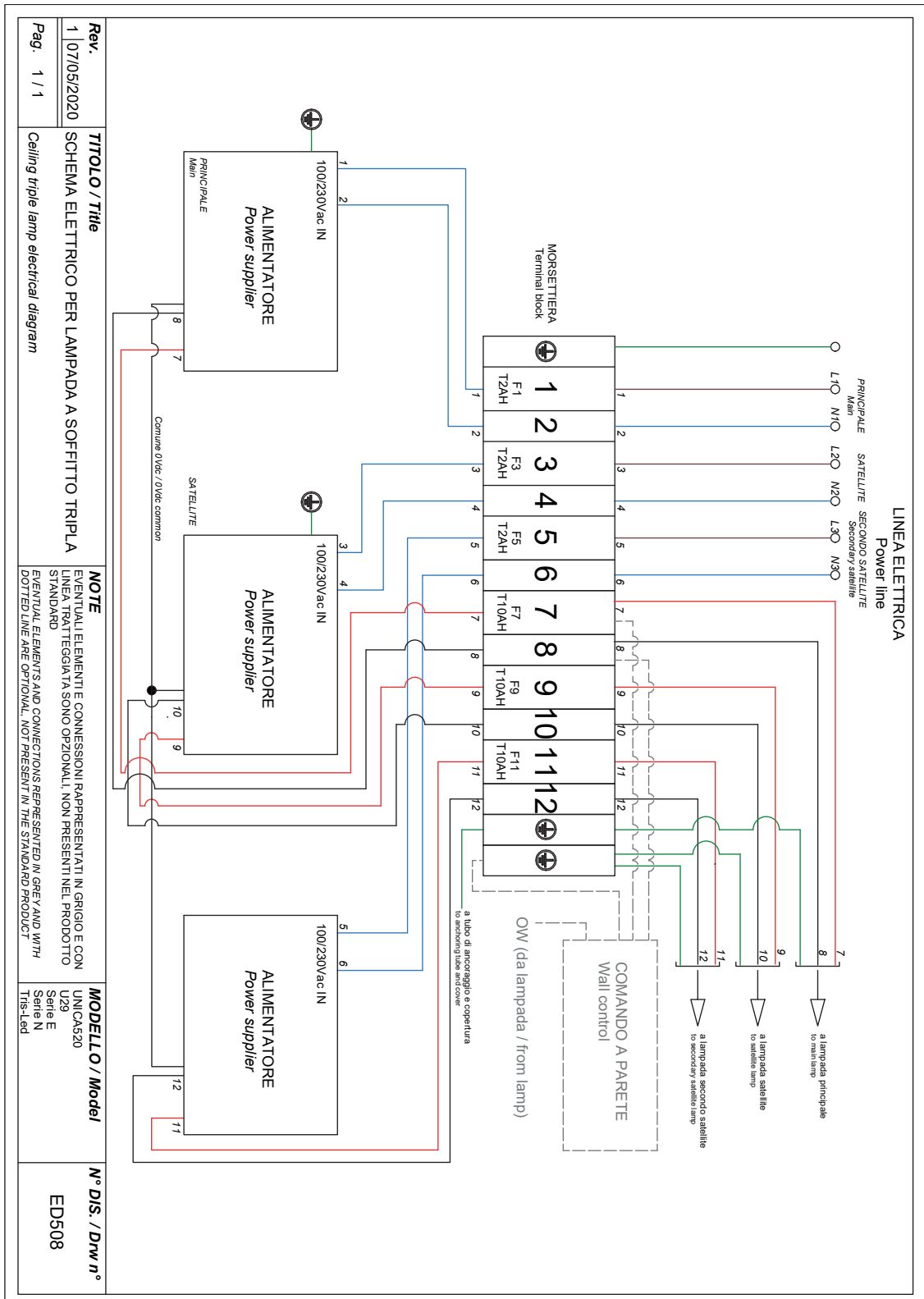
14 Electrical Schemes

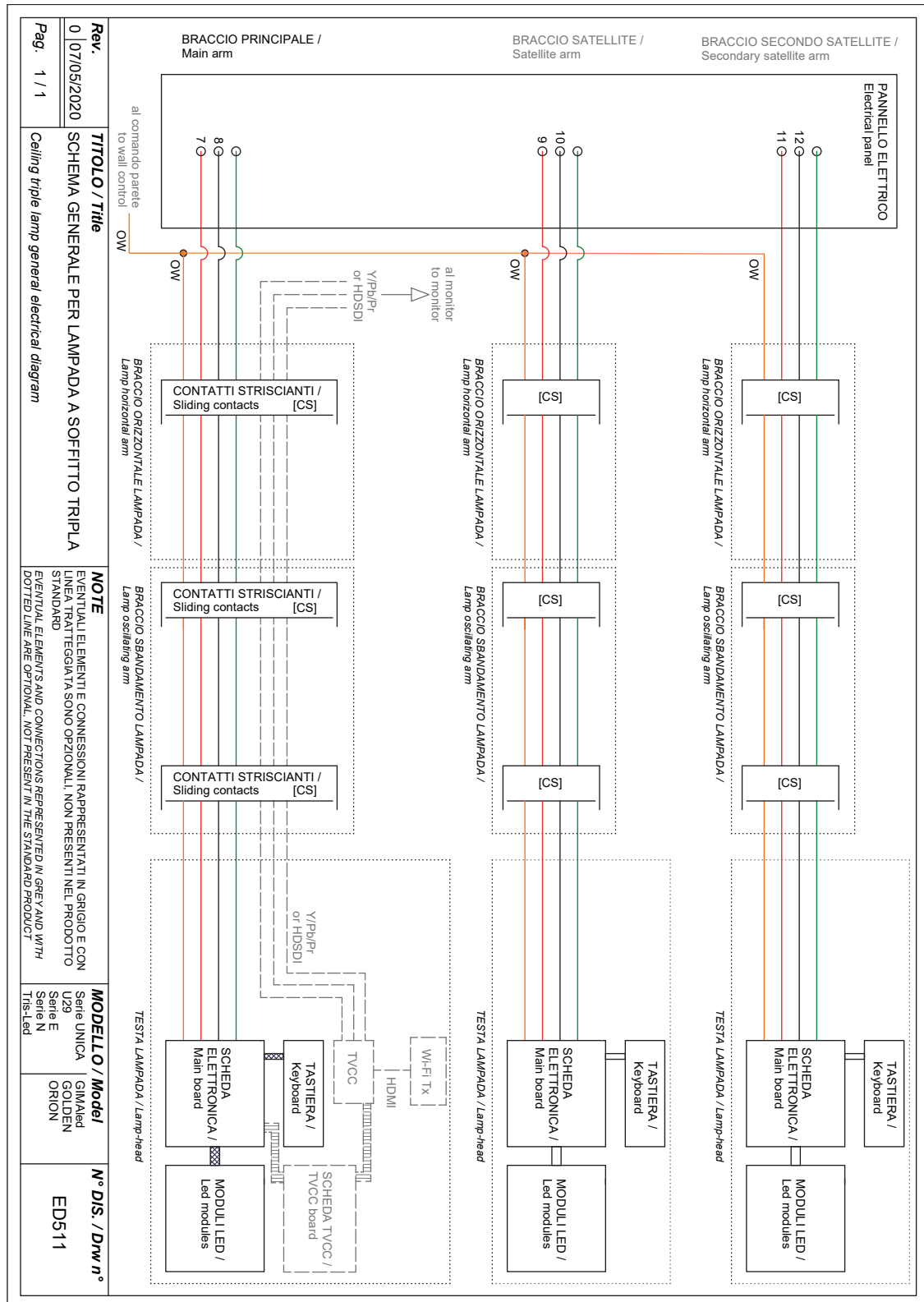




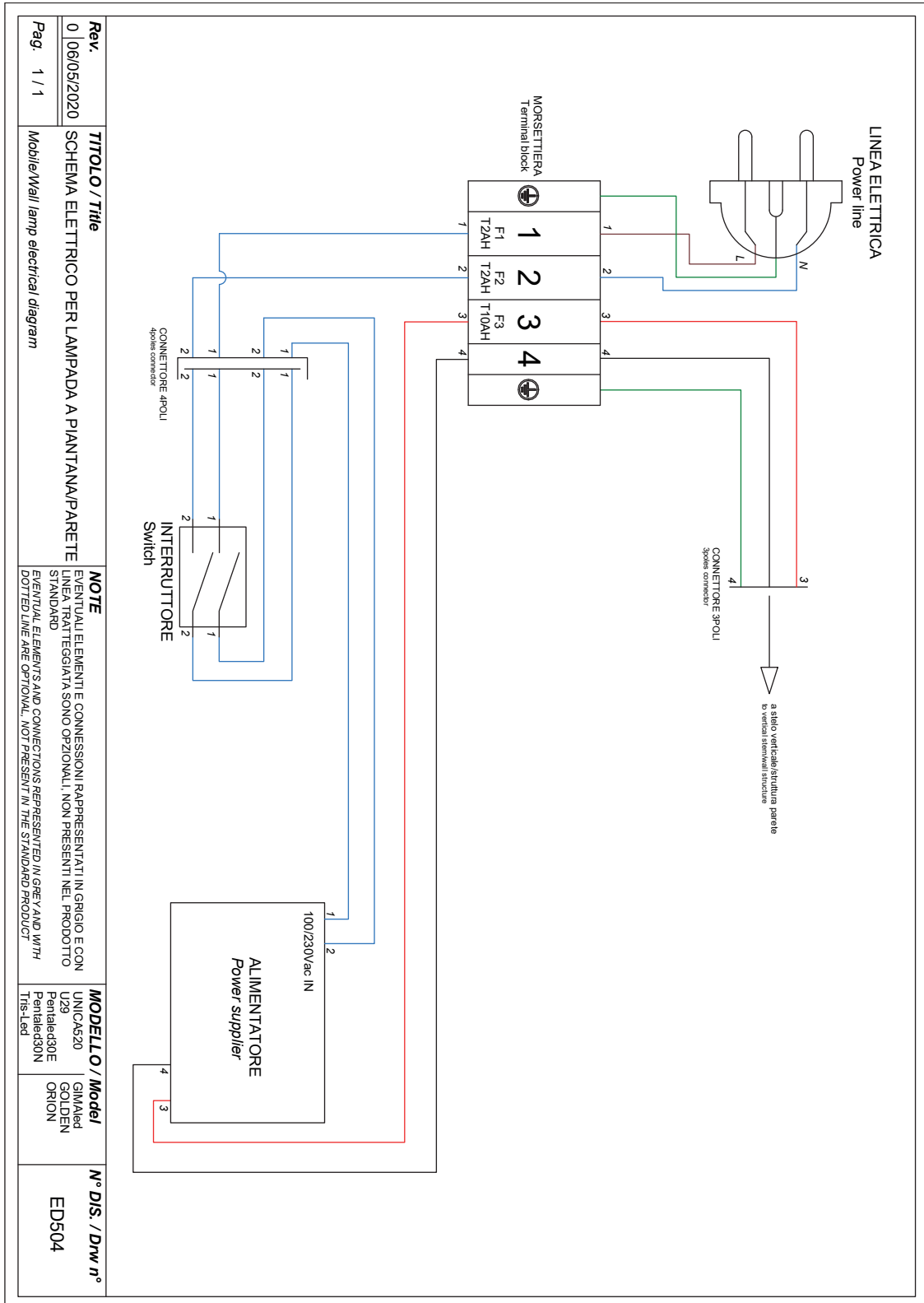


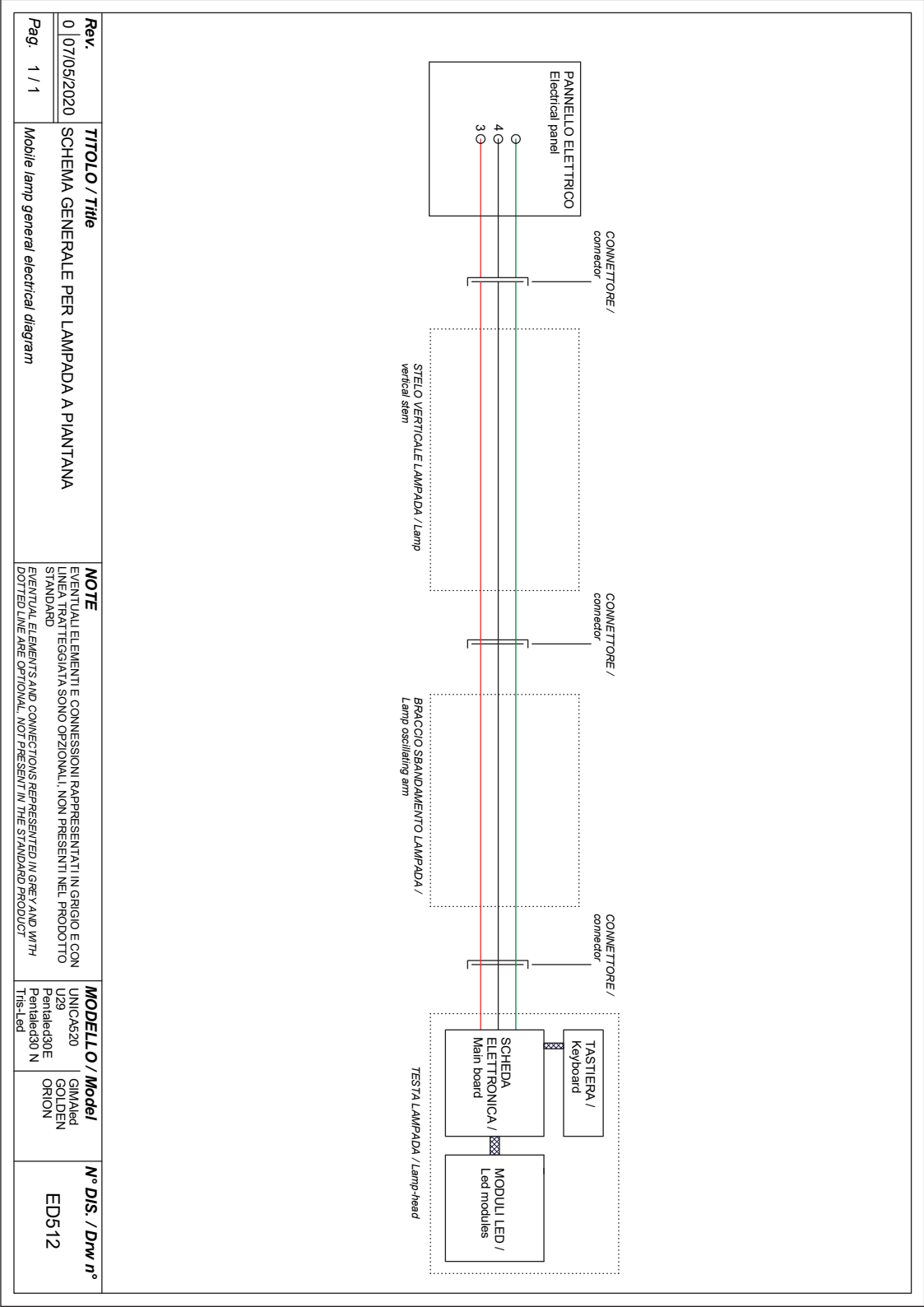


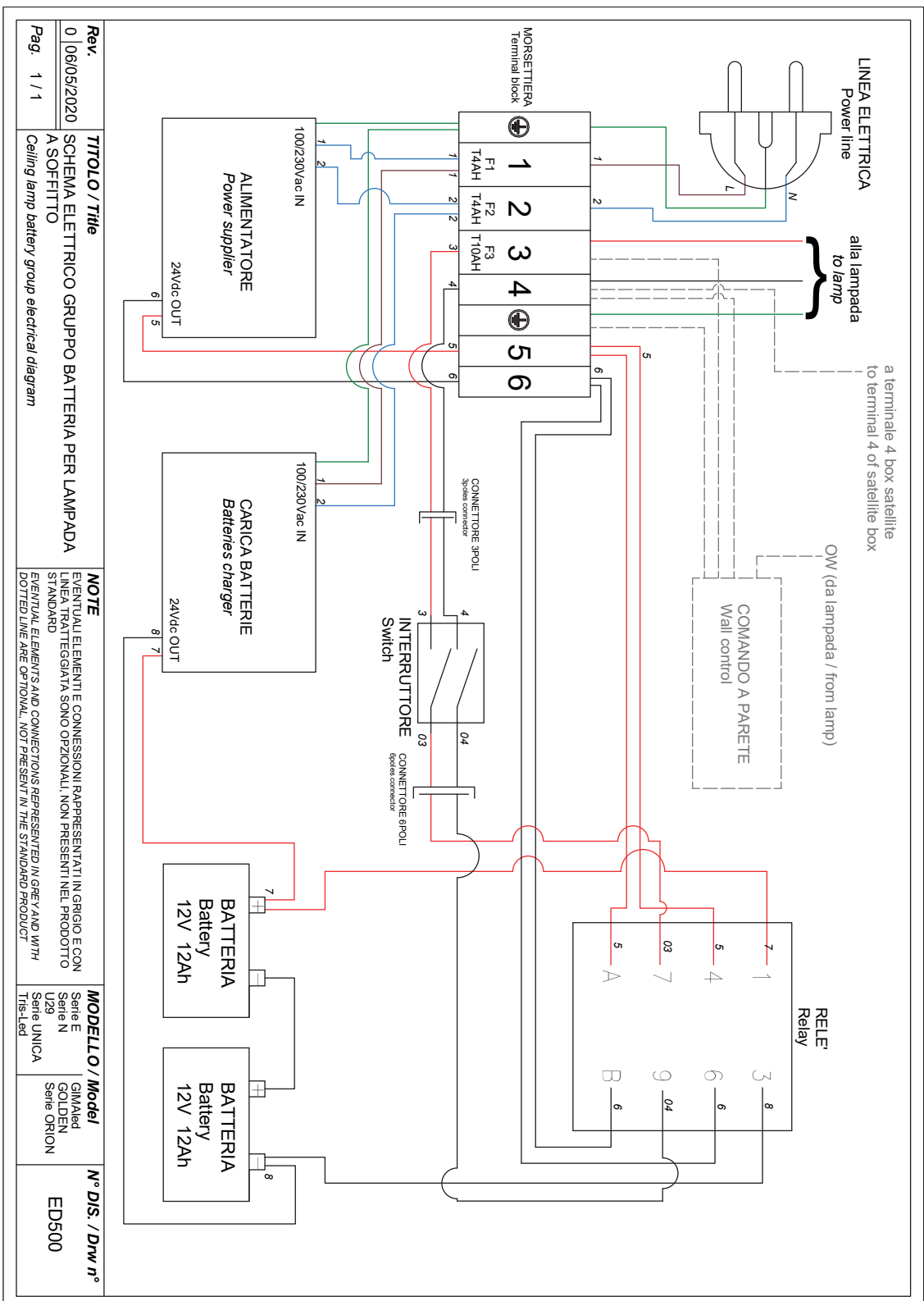


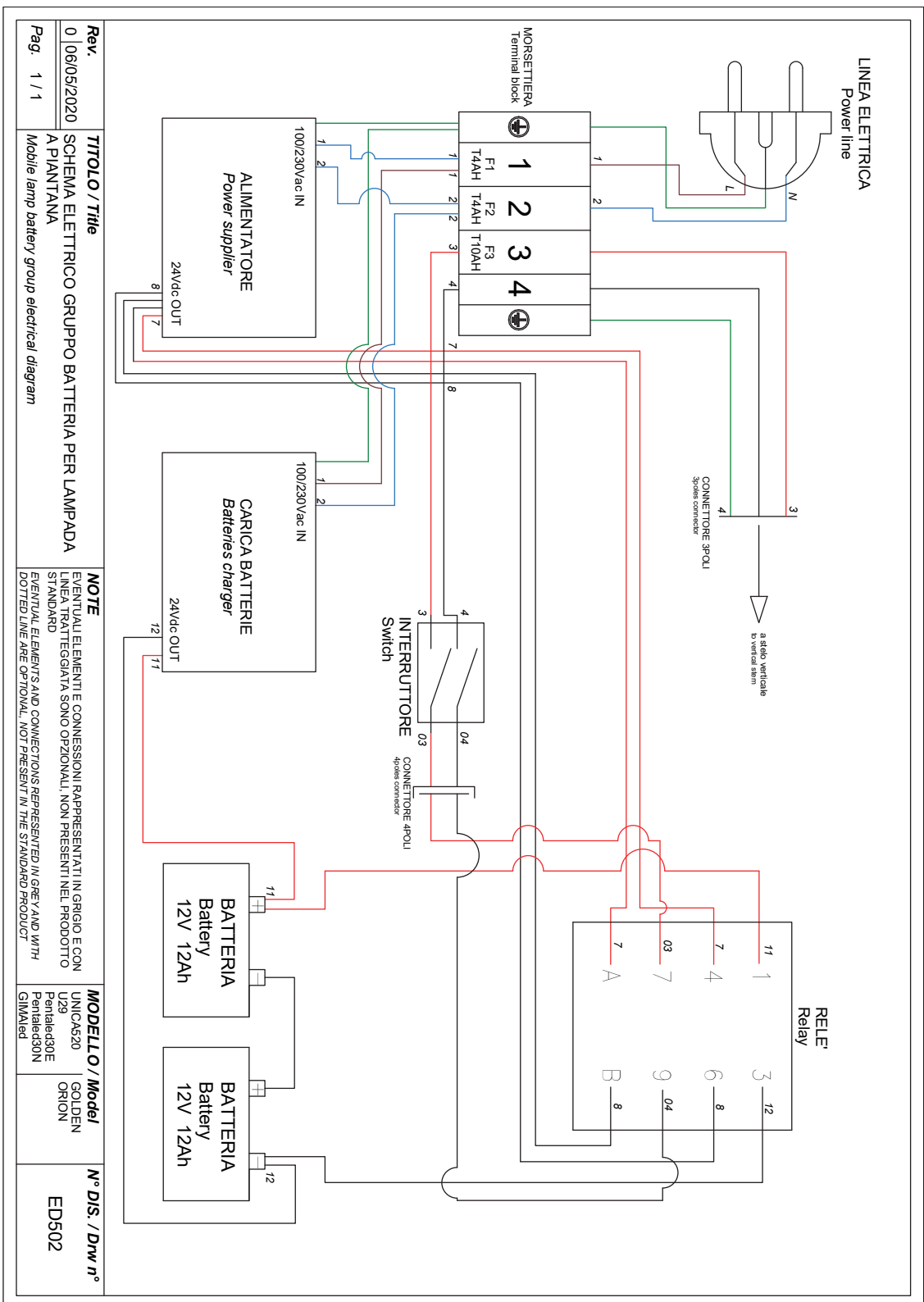


USER AND MAINTENANCE MANUAL











RIMSA P. LONGONI SRL

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