

USER GUIDE
**LOCKING SYSTEM FOR CHAIRS
SERIE PA570**

Review 2022/02



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01 INTRODUCTION













01.1 Using of manual

The manual provides using and maintenance instructions of the product as well as technical aspects, functioning, spare parts and safety.

It is recommended before the operation of the product to read carefully this manual in order to avoid damages caused by a misuse.

Do not lose this document. It should be accessible to any doubt that could appear by medical personnel. Remember that a good use and maintenance are necessary for the proper operation of the product.

01.2 Legend of Symbols

SYMBOL	EXPLANATION / DESCRIPTION
	MANUFACTURER symbol. This symbol is accompanied by the name and address of the manufacturer, adjacent to the symbol (PRODUCTOS METÁLICOS DEL BAGÉS S.L., Ctra. C-16 Km 59.5, 08650 Sallent (Barcelona)).
	Indicates the manufacturer's reference number to identify the medical device. PROMEBA, S.L. uses this symbol to set each internal reference for each configuration and business variant.
	Indicates the manufacturer's serial number to identify a specific medical device.
	Indicates the manufacturing date. The symbol must be accompanied by a manufacturing date (yyyy-mm), adjacent to the symbol.
	It is placed to inform that the product is a "Medical Device".
	CE symbol without the intervention of a Notified Organism, as a Medical Device classified as Class I according to the EU Regulation 2017/745 on Medical Devices.
	Symbol for the Unique Device Identifier.
	Symbol "See instructions for use or operating instructions".
	Symbol "caution". This symbol is placed to warn of the need for the user to refer to important precautionary information in the operating instructions, such as warnings and cautions not otherwise found on the label.
	Symbol "Caution". For a general warning.
	Warning, crushing of hands
	Indicates the temperature limits. The upper and lower temperature limits should be indicated adjacent to the horizontal lines.

01.3 Servicing request

For information of the correct interpretation of the instruction manual, the use, maintenance, installation and restoration of the product, please contact Promeba customer service: T. 93 837 12 00, email promeba@promeba.com or write to PROMEBA, S.L. - Ctra C-16 Km 59.5 · 08650 Sallent (Barcelona) · SPAIN.

01 INTRODUCTION

01.4 Demolition

When the devices are no longer suitable for use, if they have not been contaminated by any particular agent, they can be disposed of as normal solid waste, otherwise, follow the current demolition regulations

01.5 Labelling

Each product incorporates an identification label, placed on the device itself and/or on the box. It must never be removed or covered. This label includes the serial number and the product code. Please keep these numbers so you can inform the dealer if necessary.



01.6 Contraindications and adverse effects

The use of this device, if used as described in this manual, does not present any contraindications or collateral effects.

01.7 Physical requirements of the operators

Promeba locking system for stretchers is destined to professional use only.

The operators must be trained in efficient, effective and safe patient transport and must have the following minimum requirements:

- Physical capacity for operating the device
- They must be able to assess the integrity of the fixation.
- They must be able to assess any anomaly by communicating the problem to the responsible figures, interrupting the use of the devices.

It is recommended the employment of one operator equipped with strength, balance, coordination and common sense. Patient loading procedures for extremely heavy patients, operations in rough terrain and in particular situations more operators may be needed.



The capabilities of different operators should be considered before determining the role they will have in managing the device.

01.8 Intended purpose

The CHAIR LOCKS family of products is designed to anchor chairs in emergency vehicles during transport.

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01.9 General warnings

1. The product must be used by trained personnel only, having attended specific training for this device and not for similar products.
2. At least every 6 months, is important to verify if updated User Manuals are available or there is any change involving your product. This information is freely available on the website <http://promeba.com/>
3. Training routines must be registered on a special register in which the names of those trained, of the trainers, date and place are indicated. This register, which will certify the eligibility of the operators to use the Promeba, S.L. device, has to be kept for a period of 10 years after the disposal of the device itself. This register will be made available to the competent authorities and/or manufacturer if requested.
4. Promeba, S.L. is always at your disposal to plan trainings on products.
5. Before carrying out any kind of operation on the appliance (training, installation, use), the operator must carefully read the enclosed instructions, paying particular attention to the correct safety precautions and to the procedures to be followed for installation and for correct use.
6. If the instructions belong to another device and not the device received, inform the manufacturer immediately and avoid use of the device.
7. In the case of any doubts as to the correct interpretation of the instructions, please contact Promeba, S.L. for any necessary clarifications.
8. Do not allow untrained persons to help during the use of the device, because they could cause damage to the patient or to themselves.
9. Regularly check the appliance and carry out the prescribed maintenance.
10. Before each use of device the perfect operating state of the device must be checked as specified in the instruction manual. If any damage or abnormalities which could in any way influence the correct functioning and the safety of the device, of the patient and/or of the user are detected, the device must be immediately removed from service and the manufacturer must be contacted.
11. If any failure or incorrect functioning of the device is detected, it must be immediately substituted with a similar item so that the rescue procedures are guaranteed without any interruption.
12. Use of the device in anyway other than described in this manual is forbidden.
13. Do not alter or modify in any way the appliance; any such interference could cause malfunctions and injury to the patient and/or rescuer.
14. The appliance must not in any way be tampered with (modification, adjustment, addition, replacement). In such cases all responsibility will be denied for any malfunctions or injuries caused by the appliance itself; moreover CE certification and product warranty will be considered void.
15. Those who modify or have modified, prepare or have prepared medical appliances in such a way that they no longer serve the purpose for which they were intended, or no longer supply the intended service, must satisfy the valid conditions for the introduction onto the market.
16. Handle with care.
17. Ensure that all the necessary precautions are taken in order to avoid the hazards that can arise as the result of contact with blood or body fluids.

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18. Register and store with these instructions: lot number, place and date of purchase, first date of use, date of checks, name of users, any comments.
19. When the device is being used, the assistance of qualified staff must be guaranteed.
20. Do not store the device underneath any heavy objects which could cause structural damage.
21. Store in a cool, dry, dark place and do not expose to direct sun.
22. Store and transport device in its original packaging.
23. The device must not be exposed or come into contact with any source of combustion or inflammable agents.
24. Position and adjust the device taking care not to cause any obstruction to rescuers and or any other rescue equipment.
25. Attention: laboratory testing, post production tests and instruction manuals cannot always consider every possible scenario for use. This means that in some cases the performance of the product could be notable different from results to date obtained. Instructions are continually being updated and are under tight surveillance of fully qualified staffs with adequate technical formation.
26. Both public and private operators are obliged to inform the manufacturer of any measures that should be adopted to guarantee the safety and the health of the patients and the users of any medical device.
27. Promptly notify PROMEBBA regarding any revisions to be made by manufacturer in order to guarantee the conformity of the product to the territory's legal specifications (including those resulting from rules and/or norms of other nature).
28. As a distributor or end users of products manufactured and/or marketed by Promeba, S.L., you are strictly required to have a basic knowledge of any legal requirements existing in the goods final destination country applying to the devices contained in this supply (including laws and norms regarding technical specifications and/or safety requirements) and therefore you are also strictly required to have the necessary knowledge to guarantee all aspects regarding the total conformity of the products to the regulations in the relevant territory.
29. Act with all due care and diligence, and contribute to ensure conformity to general safety requirements of all devices marketed in the territory, by providing final users with all necessary information for carrying out periodical checks on their devices, as specified in the relevant user manual.
30. Actively contribute to product safety checks on products sold, by communicating any relevant risk analysis information both to the manufacturer and to any competent authorities so that the necessary actions can be promptly taken.
31. Be aware that in the event of any failure to conform to the above mentioned requirements you will be deemed fully responsible for all damages that might occur. Therefore we expressly disclaim any responsibility and/or liability for your non-compliance with the present "Regulatory provisions".

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01.10 Specific warnings

1. Establish a maintenance program and periodic testing, identifying a reference employee. The person to whom the ordinary maintenance of the device is entrusted must ensure the basic requirements foreseen by the manufacturer in the user's manual.
2. Always wear gloves during maintenance procedures.
3. Use only accessories/spare parts that are original or approved by Promeba, S.L. when carrying out any operation, to avoid causing any alteration or modification to the device, otherwise we assume no responsibility for the proper functioning or damage resulting from device to the patient or the operator and warranty will be considered void.
4. Always respect the maximum capacity of the device to be anchored in the closure.
5. The locking system must be installed in accordance with the specifications provided by the manufacturer and ensuring its compatibility with technical specifications of the chair.
6. Check that the configuration and the installation of locking system are suitable for the chair that is being used to ensure the proper use of the chair.
7. Do not use bleach to disinfect the product, neither use water jets nor pressurized air. Use a hydro alcoholic based disinfectant and wash with water.
8. The device and all its components, after washing, should be allowed to dry completely before storing. Do not use drying machines.
9. Do not lubricate any part other than the one described in this manual. Lubrication must be carried out after cleaning and complete drying.
10. Avoid contact with sharp objects.
11. Do not use the device if it has some kind of damage or poor cleaning condition.
12. Do not disassemble parts or components of the device, except those that are expressly indicated in this manual.
13. For the use of the device, one operator in suitable physical conditions is needed, with strength, balance, coordination, and common sense and must be trained on the correct functioning of the device Promeba, S.L. stretcher.
15. The laws relating to safety in the workplace must always be respected. Promeba S.L. is not responsible for damages caused by non-compliance with these requirements.
16. The installation of the locking system on the vehicle floor must be carried out ensuring compliance with the safety requirements of EN 1789. In order to be in compliance with the requirements of EN 1789, the installer has the duty to check the fixation based on the exact position on the floor and based on the structure of the floor itself.
17. The device is intended to be used on emergency vehicles.
18. The user must not have impairments that prevent proper reading and interpretation of information contained on the device labelling and in this manual, and that could prevent proper operation of controls.
19. Before and after each use, check the proper functioning of the device; if alteration or performance is noticed, it is necessary to restore its security status before the next use.

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20. If the product comes with accessories with limited lifespan, do not use them after the expiration date.
21. Do not use the device if it has not been subjected to scheduled maintenance or required for normal use.
22. Make sure that the scheduled maintenance has been carried out.
23. Promeba S.L. will not accept responsibilities in the case that the locking system is not used as described in this user's manual.
24. Do not alter or modify the locking system arbitrarily to make it fit into the ambulance: the modification may cause unforeseeable functioning and damages to the patient and operators. In any case the warranty will be lost.
25. Condensation, water, ice and accumulations of dust can affect the correct operation of the device.
26. Follow the procedures approved by the Emergency Medical Services for the immobilization and transport of patients.
27. Make sure that the product is well anchored in its corresponding anchor points.

01.11 Residual risks

The residual risks listed below have been identified only with reference to the intended use of the device:

1. Use by untrained personnel may result in injury to patient, operator, or third parties.
2. Inappropriate disinfection procedures can create a risk of cross infection.
3. Installation carried out by untrained personnel, could result in inadequate fastening of the device inside the vehicle, with consequent risks related to its instability or mobility.
4. Failure to comply with the warnings for operators can create risks.
5. Failure to read and understand the product instructions can result in injury to the patient and operators.
6. Loading of the stretcher on the ambulance could need great effort from the operator, resulting in injuries. Pay attention during loading and unloading of the stretcher and, in case of heavy weight, consider that at least 2 operators are required.

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01.12 Reference standards

REFERENCE	TITLE OF DOCUMENT
UNI EN ISO 1865-5	Patient handling equipment used in road ambulances. Part 1. Table support brancard.
UNI EN 1789	Medical vehicles and their equipment. Road ambulances



As a distributor or end user of the products manufactured and/or distributed by Promeba, S.L., you are strictly obliged to know the legal provisions in force in the country of destination of the goods, applicable to the devices to be supplied (including the regulations relating to technical specifications and/or safety requirements) and, therefore, understand the necessary requirements to ensure compliance of the products themselves with all the legal requirements of the territory.

01.13 Life span

If used as described in the following instructions, this device has a useful life of 10 years from the date of purchase.

This useful life can be extended with annual reviews carried out by the manufacturer, which uses specialized and authorized internal and external technicians.

In case these annual checks are not carried out, the device must be disposed of according to the information in paragraph 01.4 and the manufacturer must be notified.

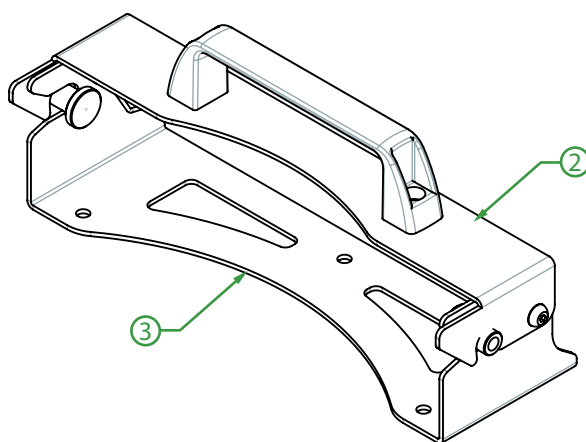
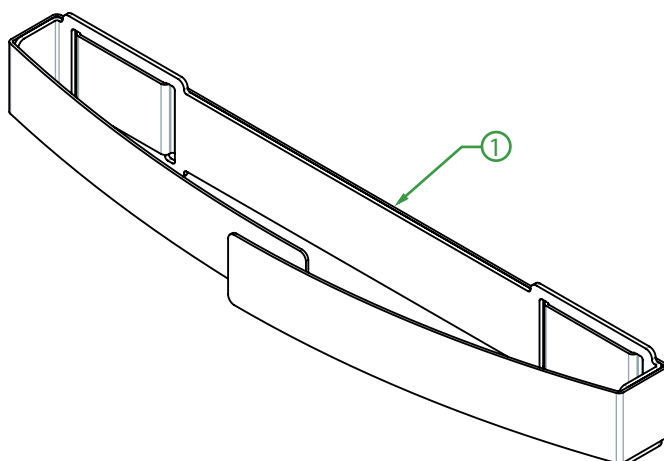
Only the manufacturer or an authorized center can extend the life of the device, if it meets the safety requirements.

Promeba, S.L. will not accept any responsibility for malfunction or damage caused by the use of devices that have not been checked by the manufacturer or authorized center, or that have exceeded the maximum allowed useful life.

02 PRODUCT DESCRIPTION

02.1 Main components

N°	DESCRIPTION OF COMPONENTS
1	Belt support
2	Upper body
3	Lower body

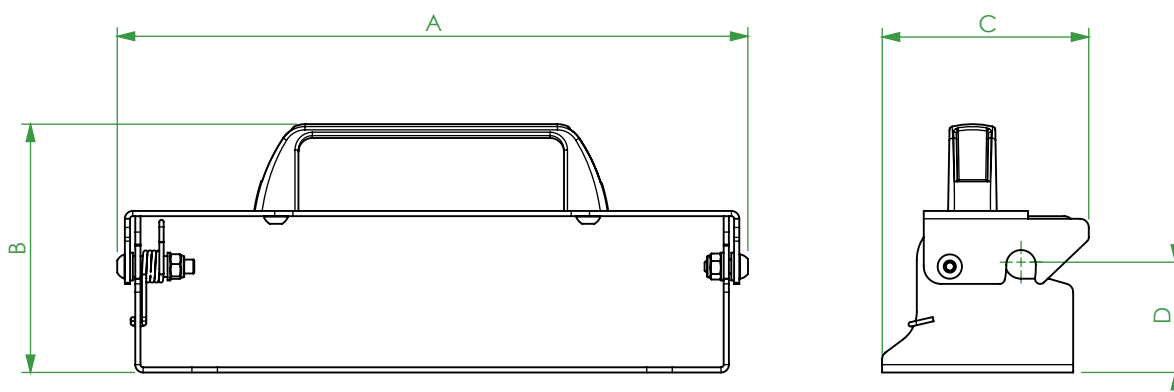


The drawing is indicative. Depending on the model obtained, it may present design variations according to its performance.

02 PRODUCT DESCRIPTION

02.2 Technical data sheet

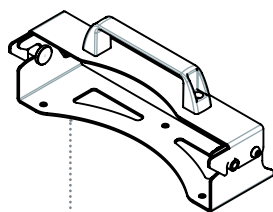
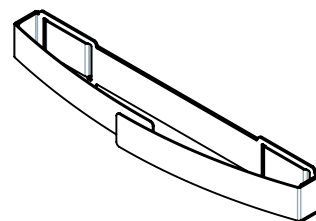
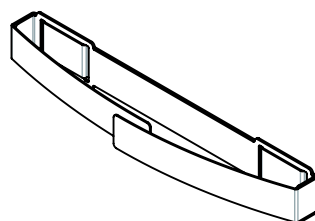
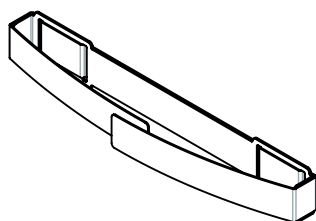
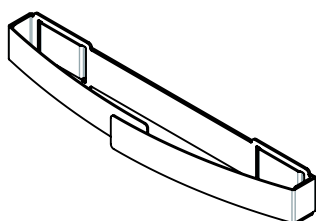
MODELS	PA570 (horizontal)	PA570 (vertical)	PA575 (horizontal)	PA575 (vertical)
Length (A)	365 ±5 mm	365 ±5 mm	365 ±5 mm	365 ±5 mm
Width (B)	120 ±5 mm	145 ±5 mm	150 ±5 mm	180 ±5 mm
Height (C)	145 ±5 mm	120 ±5 mm	180 ±5 mm	150 ±5 mm
Anchor point height (D)	64 mm	80 mm	101 mm	110mm
Weight	3,7 Kg	3,7 Kg	4,5 Kg	4,5 Kg
Recommended operators	2	2	2	2
Needed operators	1	1	1	1
Operation temperature	-20 /+40 °C	-20 /+40 °C	-20 /+40 °C	-20 /+40 °C



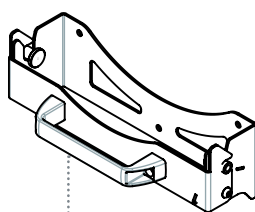
02 PRODUCT DESCRIPTION

02.3 Features

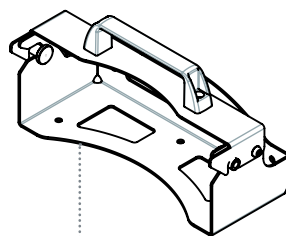
MODELS	PA570 (horizontal)	PA570 (vertical)	PA575 (horizontal)	PA575 (vertical)
Fixation system for PROMEBA chairs PS-170, PS-171 y PS-176	X	X		
Fixation system for PROMEBA chairs PS-154 y PS-155			X	
Fixation system for PROMEBA chairs PS-150 y PS-152				X
It can be installed on any flat wall or directly on the floor of the ambulance	X	X	X	X
Structure in high strength steel in painted finish	X	X	X	X
Ergonomic handle release	X	X	X	X
Belt with buckle to adjust the upper part of the chair	X	X	X	X



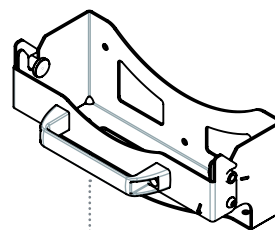
PA570 - HORIZONTAL



PA570 - VERTICAL



PA575 - HORIZONTAL



PA575 - VERTICAL

03 OPERATION

03.1 Transport and storage

Before transporting the product, make sure that it is properly packed.

During transportation, always secure the load. Transport the load leveled and following all the precepts and rules for the transport of loads, also making sure that there are no risks of bumps or falls during the transport itself.

Product damage caused during transport and handling is not covered by warranty. Repair or replacement of damaged parts is the customer's responsibility.

To unpack the product, put the box on a flat and stable surface, open the seal being careful not to damage the inside and remove the product from inside the box.

Keep the original packaging for use in any other transportation and for storage.

The device should be stored in a cool, dry place, away from direct sunlight. It must not be in contact with chemical substances or agents that can cause damage and reduce its safety performance.

During storage, do not place heavy materials on the device. The product should not be considered or used as a shelf for any type of material.

03.2 Preparation

Upon receipt of the product:

- Remove the packaging and expose the material so that all components are visible.
- Check that all components/parts on the attached list are present.

The device must be checked before each use to reveal any malfunction and/or damage caused by transport and/or storage. In particular, check:

- General functionality of the device
- Cleaning of the device (remember that the lack of cleaning can cause the spread of infections)
- Absence of cuts, holes, breaks in the structure
- Correct fixing of all nuts, bolts and screws
- State of moving parts
- Component integrity
- Lubrication of moving parts
- The state of the springs
- That the welds are intact, without cracks or breaks.
- That there are no tubes or metal sheets that have bends or cracks.
- Make sure that the vehicle in which you want to install the device is manufactured in accordance with EN 1789 standard.

If the above conditions are met, the device can be considered ready for use; otherwise, it should be immediately removed from service and contact the manufacturer.

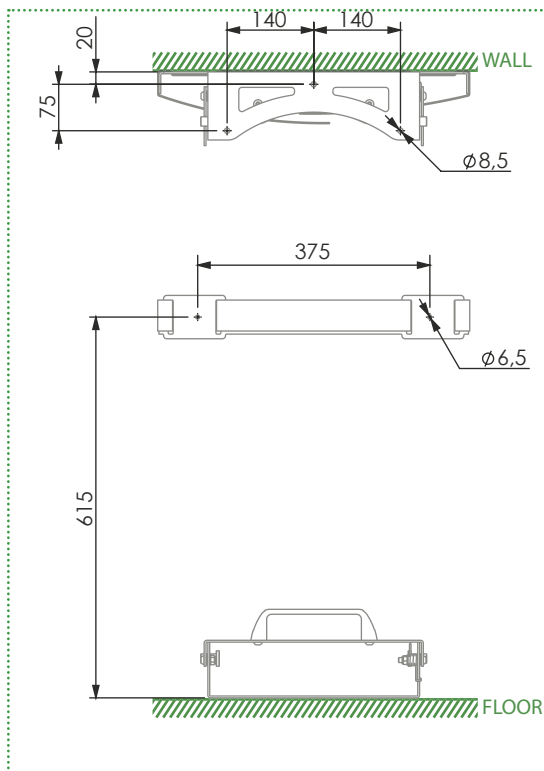
03 OPERATION

03.2.1 ANCHOR POINTS

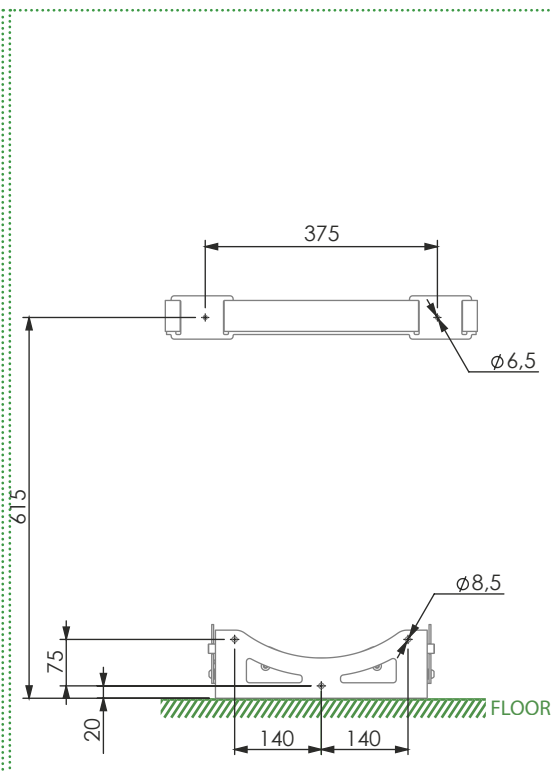
Before putting the locking system into operation, you must ensure that each and every one of the anchor points is well secured.

It is recommended to anchor the closure in the ambulance horizontally whenever possible, but in case the chair requires it, both closures also allow vertical anchoring.

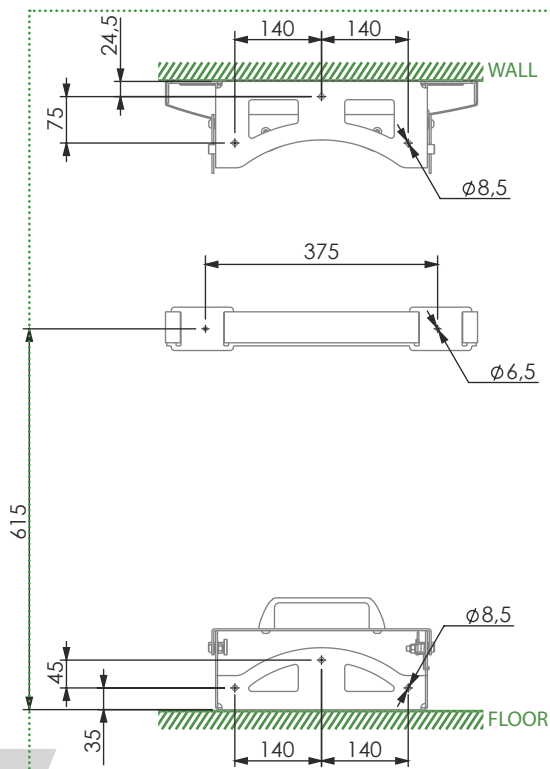
PA570 CLOSURE HORIZONTAL ANCHORAGE



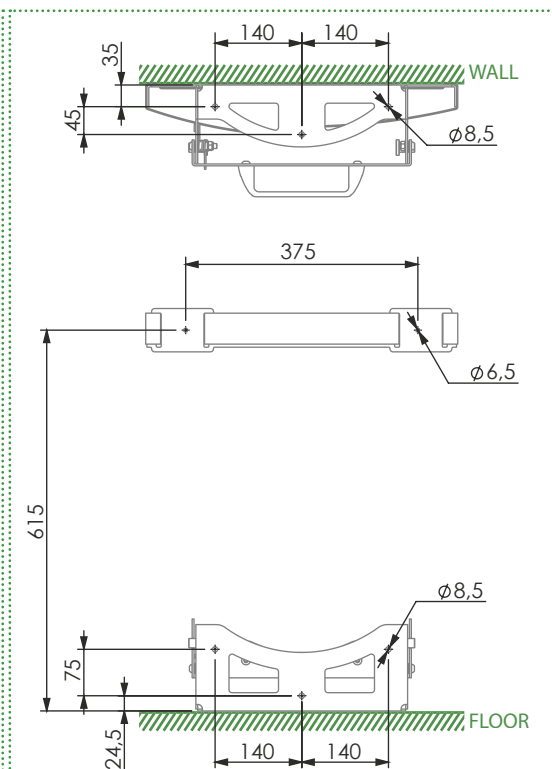
PA570 CLOSURE VERTICAL ANCHORAGE



PA575 CLOSURE HORIZONTAL ANCHORAGE



PA575 CLOSURE VERTICAL ANCHORAGE



03.3 Functioning

To place a chair in the locking system follow the next steps.

Vertically anchored closure:

LOAD:

- Fold the chair and load it into the ambulance.
- Bring the rear bar of the chair closer to the anchoring opening of the closure that is located in the upper part of it (see figure 1).
- When the bar is in position, push hard so that the lock opens and the seat bar is in the locked position.
- With the belts, adjust the upper part of the chair so that it is fixed.

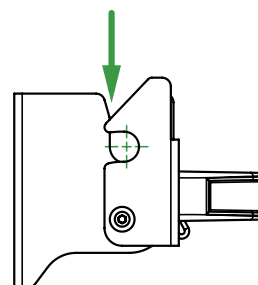


FIGURE 1

UNLOAD:

- Release the seat from the lock by pushing the handle down with your hand or foot.
- Pull the seat up, keeping the locking system open until the seat is free.

Horizontally anchored closure:

LOAD:

- Fold the chair and load it into the ambulance.
- Bring the rear bar of the chair closer to the anchoring opening of the closure located on the front of it (see figure 2).
- When the bar is in position, push hard so that the lock opens and the seat bar is in the locked position.
- With the belts, adjust the upper part of the chair so that it is fixed.

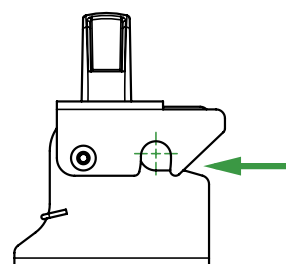


FIGURE 2

UNLOAD:

- Release the seat from the lock by pushing the handle backwards with your hand or foot.
- Pull the seat forward, keeping the locking system open until the seat is free.

04 GENERAL MAINTENANCE

CLEANING

It is essential to keep the equipment clean to ensure a proper use and durability of the set. A thorough clean must be done periodically, especially in areas exposed to dirt that may be damaged such as gears or mobile elements.

Do not use high-pressure cleaning systems, neither bleach to disinfect the product, they can damage it. Instead, use a hydroalcoholic based disinfectant, wash with water and let it dry naturally, do not use sources of direct heat to dry.

MAKE SURE TO KEEP AREAS FROM FIGURE 3 FREE OF DIRT. PAY SPECIAL CARE IN ITS CLEANING TO ENSURE PROPER FUNCTIONING OF THE ASSEMBLY.

MOBILE ELEMENTS

Due to the intense and continuous use of mobile elements such as levers or commands, periodically examine its proper operating. Inspect the mechanical connections that may exist.

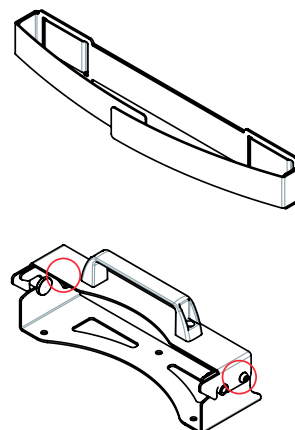


FIGURA 3

LUBRICATE

Generally, all moving parts must be lubricated. Our products leave the factory completely lubricated. However, it is possible that the elements lose lubrication with the passage of time and the use of the product, either due to loss of lubricant or dirt.

Periodically clean and lubricate affected areas according to manufacturer's specifications.

Check for loose, missing or worn parts. Periodically inspect all moving parts to ensure components are tight.

WEAR AREAS

Inspecting regularly on the system components for signs of wear is a preventive measure that can reduce breakdowns. Check possible lubricant leakages, grooves or bearing in poor condition.

MECHANICAL FIXING

We call mechanical fixing elements to the components used to fix the product as a whole, mainly screws and derivatives.

To some terms of use, due to vibrations or impacts, certain elements may lose their tightening torque or fixing properties. Periodically review that there are no loose elements, especially on moving parts. Please note and always respect the recommended tightening torques.

REPLACEMENT OF COMPONENTS

In the event that certain mechanical parts need to be replaced by qualified service personnel, they should contact our sales department for more information on ordering spare parts and their installation.

MAINTENANCE SUMMARY	EVERY USE	WHEN NEEDED	EVERY MONTH	EVERY YEAR	EVERY 4 YEARS
DISINFECT	X				
CLEAN		X			
INSPECT		X	X		
LUBRICATE		X		X	
SPRINGS REPLACEMENT				X	

05 SPARE PARTS AND ACCESSORIES

To request a spare part, indicate:

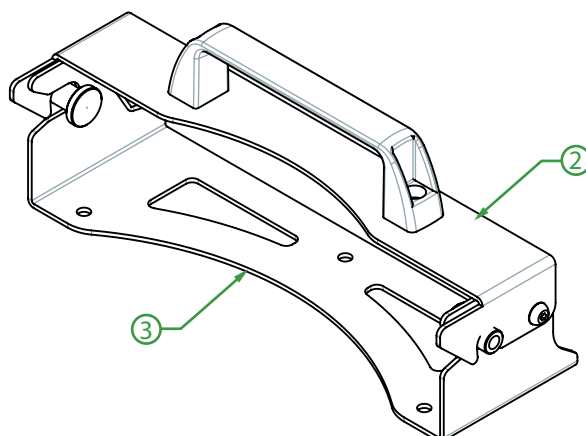
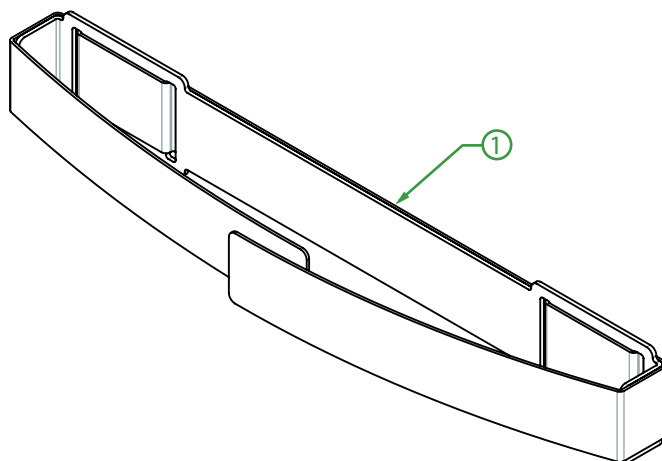
- **Model** of locking system that needs the replacement
- **Number and description** of the set in which the needed part is located (see table on the right)
- **Point out** the part in the drawing, or the area where it is located.

Following these indications, our sales department will provide you with a detailed drawing of the set so that you can specify the reference of the required spare part.



The drawing is indicative. Depending on the model obtained, it may present design variations according to its performance.

N°	DESCRIPTION OF COMPONENTS
1	Belt support
2	Upper body
3	Lower body



6 TRAINING REGISTER

The product must be used by trained personnel only, having attended specific training for this device and not for similar products.

Keep this document at least 10 years from the end of life of the device.

PLACE AND DATE	NAME OF THE OPERATOR	NAME OF TRAINER	TYPE OF TRAINING

7 MAINTENANCE REGISTER

Perform the required maintenance as indicated by the manufacturer in this user's manual.

Keep this document at least 10 years from the end of life of the device.

[illegible]

08 LEGAL NOTICES

This document may contain technical inaccuracies or typographical errors.

Changes are periodically added to the information herein; these changes will be incorporated in new editions of the publication.

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Promeba, S.L. can use or distribute the information that the client provides in whatever way they see fit, without incurring any obligation with the client.

09 PRODUCT WARRANTY

Promeba, S.L. guarantees that its products have satisfactorily passed all established quality controls, both functional and material. The duration of the guarantee is 2 years from the date of purchase of the product.

This guarantee will be valid only when it is presented with the original invoice or proof of purchase (indicating the date of purchase, model and the name of the distributor) together with the defective product during the period covered by the guarantee. Promeba, S.L. reserves the right not to offer the free warranty service if the indicated documents are not presented or if the information they contain is incomplete or illegible.

1. This warranty will not apply if the model name or serial number of the product has been altered, erased, disappeared or becomes illegible.

2. This guarantee does not cover transportation costs or risks derived from the transportation of your product to and from Promeba, S.L.

3. This warranty does not cover any of the following:

a) Periodic maintenance and repair or replacement of parts derived from normal wear and tear.

b) Expendable material (components that are expected to need periodic replacement during the life of the product, such as non-rechargeable batteries, light bulbs, etc.).

c) Damages or defects derived from the improper use, operation or treatment of the product and not due to normal use of the product.

d) Damages derived from:

i. Misuse, including:

- Treatment that results in damages or physical, superficial or appearance changes of the product.

- Installation, use or storage of the product in a way that does not respect the instructions described by Promeba, S.L.

- Maintenance of the product in a way that does not respect the instructions of Promeba, S.L. for proper maintenance.

- Installation or use of the product in a way that does not respect the technical or safety regulations of the country where it is used or installed.

iii. Use of components not provided with the product or incorrect installation of accessory parts not previously tested.

iii. States or defects of the system in which the product is used or incorporated with the exception of other products Promeba, S.L. designed for use with the product.

iv. Use of the product with accessories, peripheral units and other products of a type, condition or standards not established by Promeba, S.L.

v. The manufacturer or distributor will be solely responsible for deciding whether to send the parts for repair, or to replace the product in its entirety. In no case will operators be sent to repair or replacement of the product.

Except in the cases mentioned above, Promeba, S.L. will make no warranties in relation to product, performance, accuracy, reliability, or fitness for logic purpose of the equipment or other type. If this exception is not legal or contemplated by current law, Promeba, S.L. will limit or exclude your warranties only to the extent permitted by applicable law.

The only obligation on the part of Promeba, S.L. in connection with this warranty is to repair or replace parts subject to the terms and conditions of this warranty.

Promeba, S.L. is not responsible for loss or damage to products, this warranty or others, including economic loss or non-assessable damage; the price paid for the product; loss of profits, income, information, usufruct or use of the product or associated products or indirect, accidental or critical loss or damage.

This clause refers to whether the loss or damage is due to deterioration or inoperability of the associated product due to defects or unavailability of Promeba, S.L., which has caused downtime, loss of user time or business interruption.

In cases where the law prohibits or limits these liability exclusions, Promeba, S.L. will exclude or limit his liability only to the extent permitted by applicable law. For example, there are countries that prohibit the exclusion or limitation of damages caused by negligence, reckless negligence, willful misconduct, fraud and similar actions. The responsibility of Promeba, S.L. In this guarantee will not exceed, in any case, the price paid for the product, but if the current law allows only limitations of greater responsibilities, these will apply.



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Promeba, S.L. is to be considered not responsible for damages caused by the lack or the wrongness of the information here mentioned.*

SCAN QR CODE



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