Voyager



DO NOT DESTROY KEEP ENLCOSED KEY AND MANUAL WITH THE LIFT AT ALL TIMES.







Table of Contents

General Information	
Foreword	4
Suitability	4
Service and Support	
Manufacturer Information	5
Distributor Information	5
Definitions Used in this Manual	5
Intended Use	5
Operational Life	5
Product Identification	
How to Use this Manual	
Symbols Used	6
Safety Instructions	7
General Instructions	
Safe Working Load	
Important Safety Directions	
Shock Prevention	
Fire and Explosion Prevention	
Human and Environmental Safety Practices	
Battery and Battery Charger Safety Practices	
Homecare Environment Considerations	
Parts Designation	
Voyager Lift and Charging Station	
Hand Control	
How to Use the Voyager	
Before Approaching the Patient	
Transferring the Patient	
Emergency Lowering	
Emergency Brake	
Battery Information	
Charging the Battery	
Care and Maintenance	
Preventive Maintenance Schedule	
User Inspections	
Inspections by an Authorized Service Technician	
Cleaning the Lift	
Daily Checklist	
Strap Inspection	
Handling and Storage	
Battery Replacement	
Verification of the Charger's Power Source	
Sling Inspection and Care	
Annual Inspection	
UK Regulation	
Troubleshooting	
Labels on the Lift	
Technical Specifications	
Lift Dimensions	
Electromagnetic Compatibility	
Electromagnetic Compliance	
Electromagnetic Emissions	
Electromagnetic Immunity	
•	

General Information

Thank you for choosing the Voyager lift from Joerns / Healthcare.

Your Voyager is part of a series of quality products designed specially for home care, nursing homes and other health care uses.

We are dedicated to serving your needs and providing the best products available along with training that will bring your staff maximum benefit from every Joerns Healthcare product.

Please read this manual thoroughly. Contact us if you have any questions about the operation or maintenance of your Joerns / Healthcare product.

Foreword

Please read this manual in its entirety. It contains crucial information for the proper use and maintenance of the Voyager. It will help protect your product as well as ensure that it performs to your satisfaction.

Lifting and transferring a person always present a potential risk. Informations in this manual are safety related and must be read and understood to help prevent injuries.

If a serious incident occurs in relation to this medical device, affecting the user, or the patient then the user or patient should report the serious incident to the medical device manufacturer or the distributor. In the European Union, the user should also report the serious incident to the Competent Authority in the member state where they are located.

WARNING: Injuries can be attributed to the use of inadequate parts. Use only parts designated by Joerns Healthcare on your Voyager.

WARNING: Unauthorized changes on any Healthcare product may affect its safety. Joerns Healthcare will not be held responsible for any accidents, incidents or deficiencies of performance that occur as a result of any unauthorized changes to its products.

Tested as per standards by:



Suitability

The Voyager range of lifts are suitable for the following categories of lift within the working parameters of the lifts specified in the technical specifications:-

Category A - Wheelchair

Category B - Bed

Category C - Bath

Category D - Toilet/ shower chair

Category E - Floor

Category F - 90 degree rotation

The *Voyager* series are suitable for patients in the sitting, sitting/ recumbent and recumbent positions.

Service and Support

A service routine must be done on your Voyager by Joerns / Healthcare trained service staff. This will ensure the product remains safe and functional. See "Care and Maintenance" section.

Please contact your local Joerns / Healthcare agent for any of the following:

- more information.
- report an unexpected event or lack of performance,

General Information

- help in setting up, using or maintaining your Voyager,
- · replacement parts.

The agent can offer support and service programs to maximize the long-term safety, reliability and value of the product.

Additional copies of this manual can be purchased from your local Joerns / Healthcare agent. When ordering, include the *Instructions for Use* product number (001.08075.04 rev. 14).

Manufacturer Information

This product has been manufactured by: ArjoHuntleigh AB Hans Michelsensgatan 10 211 20 Malmö, SWEDEN

≅: +46 (0) 10-335 45 00∃: +46 (0) 413-138 76(♣): www.arjo.com

Distributor Information

UK Information

Joerns Healthcare Ltd, Drakes Broughton Business Park, Worcester Road, Drakes Broughton, Pershore, Worcestershire, WR10 2AG United Kingdom

Customer Care 2: +44(0)844 811 1158

: Info@joerns.co.uk
: www.joerns.co.uk

US Information

Joerns Healthcare Ltd, 2100 Design Road, Suite 100 Arlington, TX 76014 USA

≅: 800-826-0270
∃: 800-457-8827
≅: info@joerns.com
≩: www.joerns.com

Definitions Used in this Manual

WARNING:

Means: Failure to understand and follow these instructions may result in injury to yourself and others.

CAUTION:

Means: Failure to follow these instructions may cause damage to the product.

NOTE:

Means: This is important information regarding the correct use of the product.

Intended Use

The *Voyager* and its accessories are designed for lifting patients in a homecare setting, at nursing homes and other assisted living centres. Patient transfer must be done under the supervision of trained caregivers as per the instructions therein.

The product must only be used for the purposes stated above. It must be installed by Joerns / Healthcare authorized personnel as per local codes.

Operational Life

The equipment is designed and tested to support at minimum, a useful life of seven (7) years or 10,000 transfers, whichever comes first. It is subject to preventive maintenance as specified in the "Care and Maintenance" section in this manual.

The expected life for consumable products such as batteries, fuses, straps and cords is dependent upon the care and usage of the product concerned. Consumables must be maintained in accordance with published Instructions for Use and the "Preventive Maintenance Schedule".

Product Identification

The lift's identification number (specification, model, serial number) appears on a silver nameplate affixed on the back of the casing.

How to Use this Manual

WARNING: Do not attempt to use this product without fully understanding the information contained in this manual. A misuse of this unit may lead to a patient fall and to injuries.

Keep this manual with the lift and refer to it as needed. Make sure that all operators are regularly trained in the use of the product as per the information found therein.

General Information

Symbols Used

General Symbols	Key to Symbols
	Points out the date of manufacture and the address of the manufacturer
CE	CE marking indicating conformity with European Community harmonised legislation.
© s c us	Approved by the Canadian Standards Association.
REF	Manufacturer's catalogue number.
SN	Manufacturer's serial number.
ॐ ∏i	Refers to the Instructions for Use.
X	Waste Electrical and Electronic Equipment (WEEE) - do not dispose of this product in general household or commercial waste.
	Risk of pinching.
SWL	SWL represents the maximum load the lifter is rated for safe operation.
IP _{N1} N2	Degree of protection provided by enclosure. N ₁ : Ingress of particles, N ₂ : Ingress of water.
===	Direct current.
\sim	Alternating current.
<u></u>	Type BF applied part.
1	Operating Temperature Range.

General Symbols	Key to Symbols
(A)	Operating Humidity Range.
*	Keep dry.
MD	Indicates the product is a Medical Device according to EU Medical Device Regulation 2017/745

Charger Related Symbol	Key to Symbol
	Class II electrical equipment: term referring to electrical equipment in which protection against electric shock does not rely on basic insulation only.

Sling Related Symbols	Key to Symbols
(80°C)	Maximum washing temperature: 80 °C (176 °F), permanent cycle.
	Do not use bleach.
	Do not dry clean.
	Tumble dry low temperature.
	Do not iron.

Safety Instructions

The product must be put in service and used as per these safety instructions.

Anyone using the product must also have read and understood this manual.

If there is anything you are not sure about, contact your local Joerns / Healthcare agent.

General Instructions

Keep these safety instructions with the *Voyager* lift at all times.

CAUTION: The *Voyager* is for transferring patients only. Do not use the lift for any other purpose.

WARNING: Always place the sling around the patient as per the instructions enclosed. Failure to do so may result in injuries to you or to others.

CAUTION: Do not drop the lift, since it may cause internal damage that is not easily seen. If you suspect any damages to the lift, contact your local Joerns / Healthcare agent for servicing.

NOTE: Joerns Healthcare lifts are designed for Arjo/Easytrack rail systems, Joerns Healthcare slings and accessories.

Safe Working Load

The *Voyager* has been designed with a lifting capacity of 200 kg (440 lb).

WARNING: The *Voyager* is intended to be used for patients whose weight is within a specified safe working load. Do not attempt to lift more than the lowest weight limit indicated on the following:

- · the rail system;
- the "maximum load" label on the lift;
- · on the accessories;
- on the sling.

Surcharge of any of these elements may lead to a patient fall and to injuries.

Safety Instructions

Important Safety Directions

Always ensure that:

- The lift is installed by an authorized Joerns / Healthcare contractor or installer.
- · The product is used by trained caregivers.

WARNING: Before attempting a transfer, a clinical assessment of the patient's suitability for transfer should be carried out by a qualified health professional. A transfer conducted when it should not can degrade the patient's health condition.

- Special consideration is taken when transferring a patient who is connected to electrodes, catheters or other medical devices.
- · Violent impact during transfers is avoided.
- The sling is not damaged, torn or frayed and the straps are in good condition and properly fastened.
- The lifting steps found in this manual are followed.
- All controls and safety features are used only as per the rules specified in this manual. Never attempt to force a control or button on the lift.
- The charger is not stored in a shower, bath or other areas with high humidity.
- Any precautionary or instruction labels are read and those that cannot be easily read are replaced.
- The daily maintenance is carried out before using the lift.

WARNING: Joerns Healthcare warns of possible strangulation risks related to the lifting strap and hand control cable, and advises to take necessary precautions to prevent these.

WARNING: This product contains small parts (such as caps, see Fig. 6) that might present a severe danger to children if swallowed or inhaled.

CAUTION: Keep all components of the lift clean and dry.

CAUTION: Excessive exposure of the hand control to water (or other liquid) could cause malfunction of the device.

Shock Prevention

- Do not touch or use a lift with bare conductors or a damaged power cord. Electrically live equipment can result in serious injuries. If the lift or charger has any exposed or damaged wires, contact your local Joerns / Healthcare agent immediately.
- Do not splash or expose the device to water or moisture.
- Check nameplate for input voltage and frequency requirements. These ones differ by country. Do not attempt to use the lift in an area that has a different voltage and frequency requirement.
- Read battery and charger instructions thoroughly before using or storing them.

Fire and Explosion Prevention

- Do not place or store the battery under direct sunlight or near a heat source.
- Do not expose the batteries to flames.
 They might open, causing a chemical leak.
- Do not use the charger in the presence of flammable anaesthetic gases. The charger might generate a spark, which could cause an implosion.
- Do not short circuit the battery terminals.

Safety Instructions

Human and Environmental Safety Practices

- Should the battery casing crack and cause contents to come in contact with skin or clothing, rinse immediately with plenty of water.
- If contents come in contact with the eyes, rinse immediately with plenty of water and seek medical attention.
- Inhalation of the contents can cause respiratory irritation. Provide fresh air and medical attention.
- For recycling and disposal of the batteries, the rules as per the WEEE directive (Waste of Electronic and Electrical Equipment) as well as local laws and regulations must be followed. If not they may explode, leak and cause personal injury. When returning batteries, insulate their terminals with adhesive tape. Otherwise, the residual electricity in used batteries may cause fire or explosion.

Battery and Battery Charger Safety Practices

WARNING: Following the instructions is important for the safe use of the battery.

Only use batteries designed for use with the device. If unsure, do not use the battery. Make sure it belongs to the device by comparing the battery label with the technical specifications in the *Instructions for Use*. If battery type cannot be confirmed, call your Joerns Healthcare agent.

- Do not expose the battery or chargers to flames.
- To avoid bodily injury, do not crush, puncture, open, dismantle or otherwise mechanically interfere with the battery.
- Stop using the battery if any damage or deformation is noted.
- Be careful not to drop the battery.
- Only use the charger that has been supplied with the product.
- Do not charge the battery in an unventilated area.
- The charger must not be covered or exposed to dust.
- The charger is designed for dry areas only and for normal air humidity conditions.

Homecare Environment Considerations

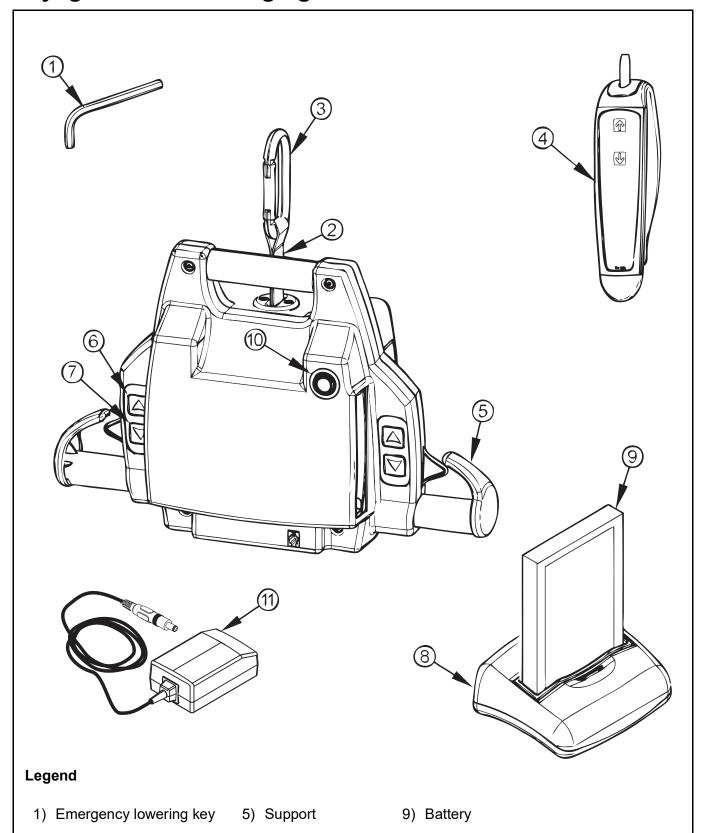
WARNING: Following the instructions is important for the safe use of the battery.

WARNING: The *Voyager* is not intended to be operated by children. Serious injuries could occur.

NOTE: Rigorous cleaning actions should be done when the *Voyager* is exposed to an animal. Pet hair trapped inside the device (through the strap opening) can reduce the product performance.

Parts Designation

Voyager Lift and Charging Station



- 2) Strap
- 3) Carabiner
- 4) Hand Control unit
- 6) UP button
- 7) DOWN button
- 8) Charging Station
- 10) Emergency stop / Battery eject button
- 11) Charger

Fig. 1

Parts Designation

Hand Control

The *Voyager*'s hand control is used to operate the lift. Each function is described in Fig. 2. The up and down buttons raise or lower the lift.

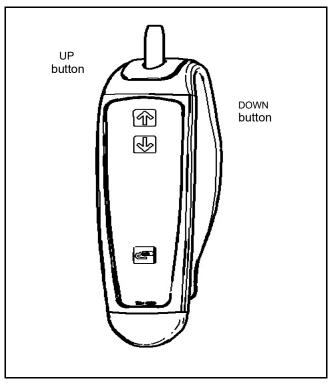


Fig. 2

WARNING: All rails must be closed with end stoppers or connected to other closed rail components. Before use, make sure all end stoppers are in place. A wrong installation of these items might lead to a patient fall and to injuries.

WARNING: The lift must never be operated by the patient. In the unlikely case of a failure, the patient might get stuck in the unit.

Before Approaching the Patient

Attendants should always tell the patient what they are about to do. Make sure to have on hand a sling that is of the correct model and of adequate size for transfer with the *Voyager*.

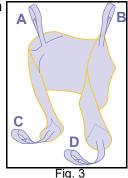
Before performing the transfer, it is important to evaluate the patient's general condition. For example, an agitated patient will require a more specific type of sling.

Attachment Method

Once the loop sling has been fitted around the patient, connect the shoulder loops of the sling to both sides of the spreader bar.

Attachments Points

The attachment points shown here are only for the purpose of the explanations below.



Cross-through method

(Legs closed with crossing straps)

This method is recommended for most general transfers.

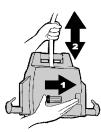


Fig. 4

Transferring the Patient

- 1) Install the sling around the patient.
- 2) Take the battery out of the charger and insert it into the lift. Make sure the silver contact points go inside the lift.
- 3) Once the patient sling is installed, unwind the strap.

There are two ways to unwind the strap:



First, you can press the down button while holding tension on the strap with the other hand (there must be tension on the strap for the lift to function).

The other way is to use the "quick-release". (1) Push the knob sideways and (2) pull the strap out of the lift.

CAUTION: Push the knob completely over to ensure that the quick-release will work properly.

The quick-release will not work if there is tension on the strap. Make sure the strap has some slack in it before attempting to slide the knob over.

DO NOT force the knob. It should slide easily.

4) Attach the carabiner to the trolley in the rail.

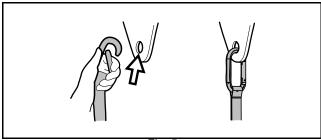
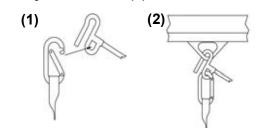


Fig. 5

5) Move the lift over to the patient.

Note: To help you reach the trolley, use the arm extension as follows:

Insert the round hook into the loop of the arm extension #A8300 (1), then hook it to the trolley in the track (2).



WARNING: Hold the lift with one hand at all times when near a patient. The proximity of the lift near the patient's face may cause the patient some discomfort.

6) Lower the lift to approximately below the patient's chin level before beginning to attach the sling straps (this is to avoid the unit from making contact with the patient's face due to the movement of the lift).

For a patient in a lying position, lower the lift near the thorax, and then install the straps.

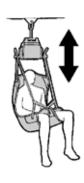
7) Attach the straps of the sling as described in the sling's documentation.

WARNING: Before lifting the patient:

- 1) Make sure that all straps are attached to the lift's supports.
- 2) Make sure the patient's arms are safely out of the way.
- Make sure the sling is not caught on any obstructions (wheelchair brake or arm of chair, etc.).

If any of the above occurs, lower the patient immediately and correct the problem.

8) Lift the patient slightly by pressing the UP control button.



- 9) Lift until the patient's buttock clears all obstacles in the transfer path, chair arm supports for example, before moving the patient. Guide the legs past any obstacle.
- 10) When the patient is located above the desired point of transfer and ready to be lowered, press the down button.

WARNING: A patient should never be left unattended during a transfer. In the unlikely event of a failure of the device, the caregiver must be ready to react.

- 11) Use the handles on the back of the sling to position the patient when transferring into a chair. Hold the handles firmly as the sling lowers and the sling will tilt back to position the patient.
- 12)Once the patient is properly seated and the straps are loose, remove the sling from the lift.

13) Slide the lift away from the patient.



14) Unhook the lift from the trolley in the rail and use the "quick release" to wind up the strap. To do so, (1) push the knob sideways and (2) hold the strap while it winds itself into the lift.

CAUTION: When using the "quick-release", hold the strap in your hand and release the knob only when the strap is not in motion. This will prevent wear of the internal components.

15) Remove the sling from around the patient.

Emergency Lowering

The *Voyager* has a safety feature that allows the lift to be lowered even if the electrical system does not operate.

Located on the right side of the lift (near the sling support), the emergency lowering device is activated by the insertion of the special key included with this manual.

To lower safely a patient, follow these steps, as illustrated in Fig. 6:

- 1) Find the rubber stopper and label on the side of the lift and pull it out. Do not discard it.
- Move the patient over a bed or chair.
- 3) Insert the 4 mm Allen key into the opening then turn it clockwise to lower the patient. The key rotation turns the motor directly, so that for each key turn, the patient will lower slightly.

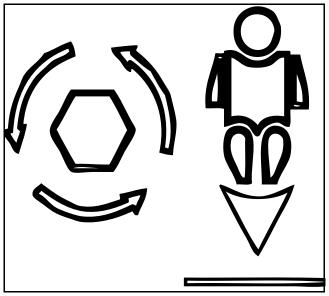


Fig. 6

CAUTION: The emergency lowering feature is to be used only in case of emergency.

4) Once the patient is lowered safely on a bed or into a wheelchair, remove the key from the lift and call a qualified technician to service your *Voyager*.

Emergency Brake

The emergency brake is an additional safety feature that automatically prevents the patient from falling in the unlikely event of a transmission or motor failure.

WARNING: After an incident has occurred and the brake was deployed, never attempt to unlock the brake or press the UP button. This may disengage the brake and make the patient fall. The emergency lowering device cannot be used.

WARNING: Never use a lift with the emergency brake deployed. Use other product to lower the patient.

The emergency brake is intended for a single activation and therefore can only be used once. Call your Joerns / Healthcare agent Service Department to arrange for the unit to be replaced.

Battery Information

For safe handling and to extend the battery lifetime, please follow and remember these instructions.

CAUTION: Not following these instructions can cause short battery life.

Only use batteries designed and labelled for use with the device. When not sure, do not use the battery. Make sure the battery belongs to the device by comparing the battery label with the technical specifications in this manuel. If battery type cannot be confirmed, call your Joerns / Healthcare agent.

Battery life depends on many factors: frequency of use, frequency of charging, temperature of operation, storage and storage time.

Make sure to have a replacement battery ready when needed. Have the facility department keep one in stock.

The battery included in the *Voyager* is 24 VDC, 2.0 Ah and rechargeable. It provides 10 to 30 transfers per battery charge.

NOTE: To prolong battery life, place the battery in the charger whenever the lift is not in use. If the low battery indicator beeps, be sure to recharge the battery as soon as possible. Charge the battery until the charging indicator light is a solid yellow before using the lift again. This will extend the life of the battery.

NOTE: Joerns / Healthcare uses sealed leadacid batteries. These batteries are not affected by any memory effect. Therefore, batteries should not be completely drained before recharging.

Charging the Battery

WARNING: Do not operate the charger with a damaged cord or if the unit has been dropped or damaged.

Do not bend the power cord by force, or place a heavy object on it. This will damage the cord and may cause fire or electrical shock.

CAUTION: Only use chargers that are specified by Joerns / Healthcare to recharge the batteries.

The steps for recharging the battery are as follows:

1) Eject the battery from the lift unit by pressing on the battery eject button.

NOTE: To avoid dropping the battery, keep your hand near the battery opening when pressing on the battery eject button.

2) Check to make sure that the green light on the charger is illuminated. Insert the battery into the charger, ensuring that the metal contact points on the battery touch the charger's contacts.

The charging indicator light (yellow) will flash while the battery is charging. When the battery is fully charged, the charging indicator light will become solid.

If the battery is low, it will take approximately two hours for a full charge.

If the battery is fully drained it could take up to 6 hours to recharge.

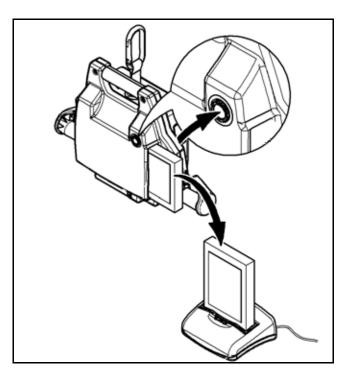


Fig. 7

NOTE: Whenever possible, leave the battery in the charger when the lift is not in use.

At a minimum, charge the battery until the charging light turns solid yellow before using again. This will extend the life of the battery.

When the lift is needed again, pull the battery out of the charger and gently slide it back into the *Voyager* lift. The metal contacts should be inside the lift.

The charger's power cord can remain plugged into the wall indefinitely, convenient for overnight recharging.

The battery can be left in the charger for an extended period of time without damaging either the charger or the battery.

If the green light does not go on when the charger is plugged into an electrical outlet, see the "Troubleshooting" section.

CAUTION: Do not attempt to use a battery not authorized by the manufacturer. The *Voyager* and its charging system use specially designed batteries. Using unauthorized batteries may seriously damage the lift or the charger.

CAUTION: The lift should not remain stored for long period of time without charging the battery. Batteries should be recharged at least every two weeks to maximize their lifespan.

Preventive Maintenance Schedule

The product is subjected to wear and tear, and the following maintenance instructions must be acted upon when specified to ensure that it remains within its original manufacturing specifications. Care and maintenance must be carried out as per the preventive maintenance schedule below.

Customer obligations must be carried out by qualified personnel as per the instructions in this manual.

WARNING: The maintenance described in the following checklist is the minimum that the manufacturer recommends. In some cases more frequent inspections should be carried out. Continuing to use this product without conducting regular inspections or when a fault is found will seriously compromise the safety of the user and of the patient. Local regulations and standards may be higher than those of the manufacturer. A load test is recommended. Service and preventive maintenance can be arranged with the manufacturer. Preventive maintenance specified in this manual can prevent accidents and reduce repair costs.

WARNING: Safety related maintenance and authorized service must be carried out by qualified personnel, fully trained in servicing procedures by Joerns / Healthcare, and equipped with correct tools and proper documentation, including Parts List and Service Manual. Failure to meet these requirements could result in personal injuries and/or unsafe product.

WARNING: Never proceed to maintenance or service while lift is in use with a patient.

User Inspections

Action/Check	INITIALLY	BEFORE EVERY USE	EVERY 2 MONTHS	EVERY 4 MONTHS	EVERY YEAR	EVERY 2 YEARS
Lift device						
INSPECT FOR MISSING HARDWARE OR BROKEN ENCLOSURE.	Х	Х			Х	
INSPECT STRAP FOR WEAR.		Х	Х			
RECHARGING BATTERY.		Х				
INSPECT THE SLING SUPPORTS ON THE LIFT FOR DAMAGE OR CRACKS.					Х	
INSPECT WHEELS IN RAIL FOR DAMAGE, RUST OR CRACKS. REPLACE IF DAMAGED.					Х	
CLEAN THE RAIL.				Х		
OVERALL INSPECTION BY AUTHORIZED PERSONNEL.					Х	
INSPECT CARABINER FOR DAMAGE OR CRACKS					Х	
VERIFY EMERGENCY STOP BUTTON.				Х		
VERIFY EMERGENCY LOWERING DEVICE.				Х		

Action/Check	INITIALLY	BEFORE EVERY USE	EVERY 2 MONTHS	EVERY 4 MONTHS	EVERY YEAR	EVERY 2 YEARS
Sling and hardware						
CHECK ALL SLING ATTACHMENTS FOR SIGN OF WEAR.		Х				
INSPECT SLING MATERIAL FOR WEAR OR DETERIORATION.		Х				
INSPECT SLING STRAPS FOR WEAR.		Х				
INSPECT FOR ANY DEFECTS OR LOOSE THREADS IN THE "STITCHED AREAS".		Х				
CLEANING SLING AS INDICATED ON THE TAG.			WHEN NEO	CESSARY		

Inspections by an Authorized Service Technician

Action/Check	INITIALLY	BEFORE EVERY USE	EVERY 2 MONTHS	EVERY 4 MONTHS	EVERY YEAR	EVERY 2 YEARS
Authorized Service						
Replace strap.						Х
Inspect the welding aspect on the frame to detect cracks.					Х	
Inspect transmission.					Х	
Inspect connecting joints for proper attachment.					Х	
Verify that the emergency brake on the drum is turning freely.					Х	
Verify emergency devices for good functioning.					Х	
Load test with SWL (maximum working capacity).					Х	

NOTE: If the product does not work as intended, immediately contact your local Joerns \prime Healthcare agent for support.

WARNING: Always reinstall the rail end stoppers (if removed) after servicing.

WARNING: Make sure the trolley is compatible with the stoppers and rail system.

Cleaning the Lift

NOTE: It is recommended to clean the lift and its accessories between each patient use.

Removing visible residues:

- 1) Use a cloth soaked with water.
- 2) Remove visible residues from the lift and its accessories (sling, handset, strap) from top to bottom.

Cleaning:

- 1) Use a damp cloth with warm water mixed with a mild detergent.
- 2) Scrub the equipment from top to bottom, removing all visible residues.
- 3) Rinse with clean water any remaining residual detergent. Then wipe clean with a dry cloth.

NOTE: Pay special attention to areas pointed in. These are most likely to enclose germs. Use a smaller brush and/or cotton swab to reach them.

CAUTION: Do not drench the product, as this could damage electrical components and cause internal corrosion.

If a hot air dryer is used to dry the lift, the temperature must not exceed 80°C (176°F).

Do not use petroleum-based solvents, as this may damage plastic parts.

Voyager's Special Areas to Clean

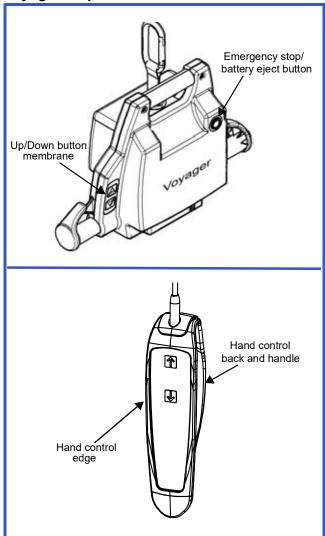


Fig. 8

To ensure a better rolling surface for the trolley wheels, clean the inside of the rail every 4 months. To do so, insert a damp cloth in the opening and slide it from one end of the rail to the other.

Daily Checklist

The following procedures must be followed before each use:

- Charge the battery.
- Inspect the lift for any damage. If the lift casing does not look properly aligned, or there are any cracks or other damage on the lift, or there are parts missing, do not use it. Contact your Joerns / Healthcare agent to have the lift serviced.
- Inspect the strap for any visible signs of wear, frays, loose threads or other damage (see Fig. 9). If there is any evidence of damage, do not use it. Contact your local Joerns / Healthcare agent to have the lift serviced.
- Inspect the sling for tears, frayed straps or loose stitching. If the sling has any of the above damage, do not use it. Contact your local Joerns / Healthcare agent to have the sling replaced or repaired.
- Inspect the carabiner at the top of the strap to ensure that it is properly attached.

WARNING: Before each use, make sure all end stoppers on the rail are in place.

NOTE: The lift cassette and the sling should be cleaned between use of different patients.

WARNING: Do not splash, drench or immerse the unit in water.

Strap Inspection

If the strap is damaged or shows signs of wear or discolouration, the acceptable load on the strap before rupture can drop rapidly and present a danger for the patient or caregiver. Joerns / Healthcare recommends a thorough inspection of the straps every two months as follows:

- Completely unwind the strap.
- 2) Look for any signs of wear or discolouration (see Fig. 9).

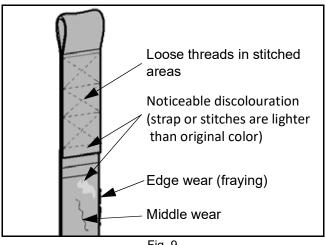


Fig. 9

WARNING: The strap must be changed if it shows any sign of wear as indicated above or any other visual defects.

The manufacturer recommends changing the strap at least every 2 years or every 2850 cycles—which ever comes first.

By continuing to use the lift without changing a damaged strap, the safety of the caregiver or patient is greatly compromised.

Handling and Storage

Avoid violent impacts while transporting the lift.

The lift should not remain stored for long periods of time without recharging the battery.

NOTE: Even if the lift is not used. Joerns / Healthcare recommends charging battery at least every two weeks. This will prevent premature aging of battery.

If you store or ship the Voyager, ensure that the power is turned off (push the battery eject button) beforehand.

Battery Replacement

Replace the battery when there is a noticeable reduction in the number of transfers that can be performed between charges. If you hear the Voyager lift beeping, see the instructions in the "Troubleshooting" section of this manual to determine if there is a problem with the battery.

For battery replacement, contact your local Joerns / Healthcare agent.

CAUTION: Do not attempt to use a battery that was not supplied by Joerns / Healthcare. The battery has been specially designed for Joerns / Healthcare charging systems. Attempting to use an unauthorized battery may seriously damage the lift and/or the charger.

Verification of the Charger's Power Source

If the light does not illuminate when the battery is correctly installed in the *Voyager* charger, try the following:

- 1) Make sure that the charger is correctly plugged into the AC outlet, and that the green light is on.
- 2) Make sure that the battery is properly inserted in the charger.
- 3) Check the power of the AC outlet on the wall.
- 4) If the charger's yellow or green light does not light up, contact your local Joerns / Healthcare agent for assistance.

Sling Inspection and Care

Refer to documentation included with slings.

WARNING: The slings should be checked before and after use and, if necessary, washed as per instructions on the sling. This is especially important when using the same equipment for another patient. This minimizes the risk of cross infection.

Annual Inspection

The *Voyager* and its accessories must be inspected annually by a qualified technician.

WARNING: The *Voyager* and accessories must be serviced every 12 months as a minimum requirement. Do not attempt to do the inspection unless you are qualified to do so. Not servicing the product may lead to patient fall, causing injuries.

UK Regulation

Note: In the UK the *Voyager* must be inspected and maintained in compliance with 1998 no 2307 health & safety: lifting operations and lifting equipment regulations 1998. Your dealer will provide you with more information.

Troubleshooting

PROBLEM	VERIFY
The unit does not work when you press the UP or DOWN buttons on the hand control.	 Make sure the hand control is plugged correctly into the unit. Use the control switches directly on the unit. If they are working, it indicates a problem with the hand control; you would then need to change the hand control.
The unit does not work when you press the UP or DOWN buttons on both sides of the lift.	 Is the battery charged and correctly installed? Ensure the battery is inserted into the lift with the metal contact points inside the lift. Place the battery into the charger and check the light on the charger. If the yellow light keeps flashing, leave the battery in the charger for approximately 2 hours. Is the "quick release" knob completely to the left (locked) position? If not, the lift will not operate. Tighten the tension on the strap to help move the "quick release" knob. Is there tension on the strap? The "anti-crush" feature will prevent the lift from working if there is no tension. If all of the previous points have been checked without success, you probably have a motor or electrical problem. You need to call your Joerns / Healthcare agent.
The unit starts and stops repetitively.	 If the load is more than safe working load, the unit will not work due to an overload protection on the motor. Has the lift been making a "beeping" sound? The battery is low and needs to be charged.
The unit emits a "beep" during use. The unit may stop lifting but you can still lower it.	Battery is low. Recharge it.
Battery is always dead after only a few transfers (3 to 5).	Replace battery with a new one, the life of the current one is probably nearly over. (See "Battery Information" in the "Care and Maintenance" section).
The light on the charger shows a solid yellow when the battery is inserted, yet the lift will only do one or two transfers.	Battery has either been dropped or damaged in some other way. Replace the battery.
The charger's green light is not illuminated.	 Is the charger plugged into a standard outlet? Try plugging the charger into another outlet to see if it will work. If the green light will still not illuminate, contact your local agent.

Troubleshooting

PROBLEM	VERIFY
When battery is inserted into the charger, the yellow light does not flash or illuminate solid.	 Check the orientation of the battery. The contact points should be inside the charger. If the yellow charge light will still not illuminate, contact your local agent.
The battery does not slide into the lift.	 Check if there is any damage to the battery that would interfere in the lift battery's slot. IF THERE IS DAMAGE - DO NOT USE THE BATTERY. Try to slide in another battery to see if there is any damage in the lift battery's slot. If so, call your Joerns / Healthcare agent. If not, discard the damaged battery.
Cannot remove battery from the lift.	 See "How to Use the Voyager" section, under "Charging the Battery" on how to correctly remove the battery. If the battery will not come out of the lift, tilt the lift to the side so that the battery opening is under the lift. With your hand guarding the opening for the battery, press the eject button on the front of the Voyager while gently tapping the lift to help jolt the battery out of the lift. ONCE THE BATTERY IS REMOVED – DISCARD BATTERY AND REPLACE. If you cannot remove the battery after trying the above, contact your local agent for assistance.

Labels on the Lift

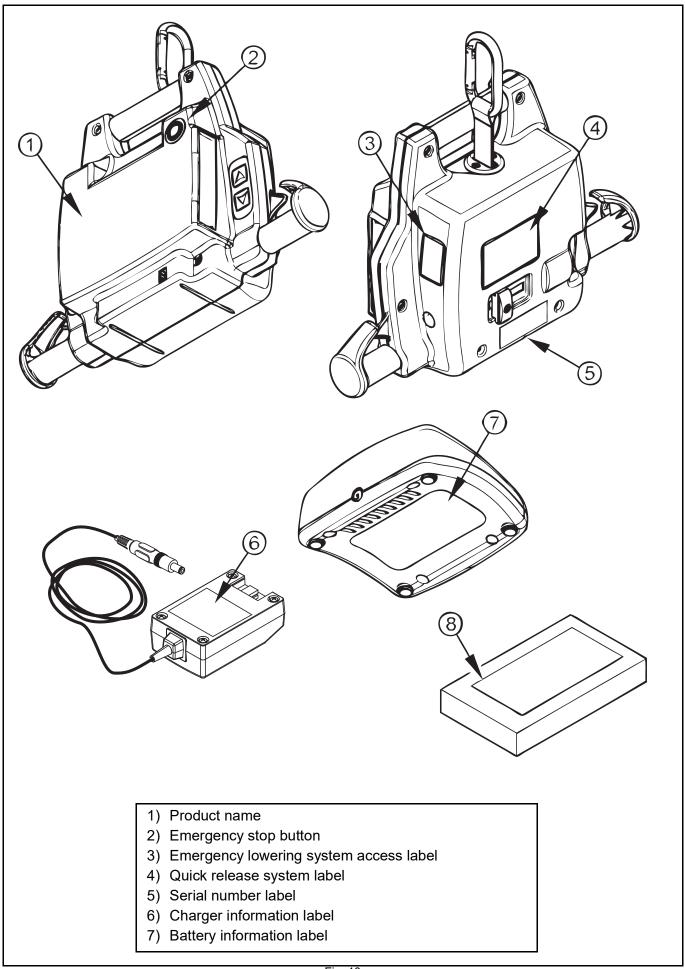


Fig. 10

Technical Specifications

PRODUCT INFORMATION			
Weight, complete	5.45 kg (12 lb)		
Lifting capacity	200 kg (440 lb)		
Strap length	2200 mm (88 in)		
Lifting speed	4.3 cm/s (1.7 in/s)		
Operating force of control	< 5 N		
ELECTRICAL			
Duty cycle	Max 10%, 6 min/hr, (max. 1 min. continuous)		
Rating	24 VDC, 10 A max.		
Noise level for either raising or lowering, with or without load	70 dBA max.		
Medical equipment	Type BF protection against electrical shock as per IEC 60601-1		
	-1 series including applicable collateral standards and national vager is compliant to ISO 10535 standard.		
cables and external antennas) show including cables specified by the equipment could result. See the	cations equipment (including peripherals such as antenna uld be used no closer than 30 cm to any part of the <i>Voyager</i> , manufacturer. Otherwise, performance degradation of this "Electromagnetic Compatibility" section for more details.		
BATTERY AND CHARGER UNIT			
Battery	Qty: 1 x rechargeable 24 VDC, 2.0 Ah		
Battery capacity	Provides up to 30 transfers with a load of 80 kg (176 lb)		
Degree of protection - Hand Control	IPX7		
Degree of protection - Voyager	IPX0 - Keep dry		
Lift - protection class - shock prevention	Internally powered equipment		
Battery Charger input	100-240 VAC, 50-60 Hz, 50 VA		
Battery Charger output	24 VDC, 1 A, 24 VA		
Battery Charger safety protection	Class 2, double insulated		
OPERATION CONDITIONS			
Ambient temperature range	+5 °C to +40 °C (+41 °F to +104 °F)		
Relative humidity range	15% to 93%, non-condensing		
Atmospheric pressure range	700 hPa to 1060 hPa (max. 2000 m)		
STORAGE CONDITIONS			
Ambient temperature range	-25 °C to +70 °C (-13 °F to +158 °F)		
Relative humidity range	< 94%, non-condensing		
Atmospheric pressure range	500 hPa to 1060 hPa		
air or oxygen, or with nitrous oxid	able in the presence of flammable anaesthetic mixtures with de. Using the <i>Voyager</i> in this environment might lead to an toreate some spark internally and ignite the gas.		
End of Life Disposal			
Battery	All batteries in the product must be recycled separately. Batteries are to be disposed in accordance with national or local regulations.		
Slings	Slings including stiffeners/stabilizers, padding material, any other textiles or polymers or plastic materials etc. should be sorted as combustible waste.		

Technical Specifications

Electrical and electronic components	Lift systems having electrical and electronic components or an electrical cord should be disassembled and recycled per Waste of Electrical and Electronic Equipment (WEEE) or in accordance with local or national regulation.
The product	Components that are primarily be made up of different kinds of metal (containing more than 90% metal by weight) for example trolleys, sling bars, rails, upright supports, etc., should be recycled as metals.

Lift Dimensions

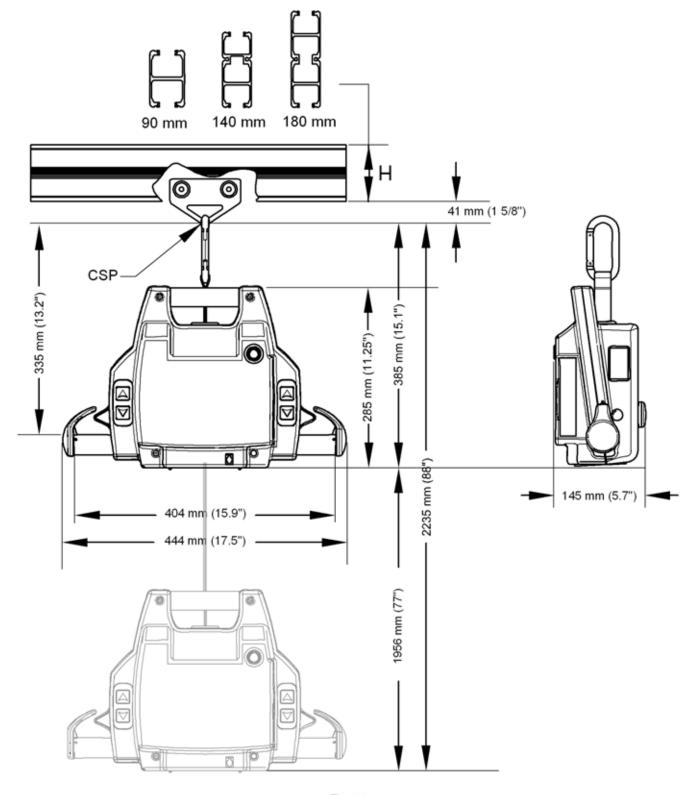


Fig. 11

Electromagnetic Compatibility

Electromagnetic Compliance

The *Voyager* has been tested for compliance with current regulatory standards regarding its capacity to block EMI (electromagnetic interference) from external sources.

Nonetheless, some procedures can help reduce electromagnetic interferences:

- Ensure that other devices in patient-monitoring and/or life-support areas comply to accepted emissions standards.
- Maximize the distance between electro-medical devices. High-powered devices may produce EMI that can affect the ceiling lift.

WARNING: Use of accessories, cables and spare parts other than those specified or provided by Arjo could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.

WARNING: Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.

WARNING: The equipment may cause radio interference or may disrupt the operation of nearby equipment. It may be necessary to take action, such as reorienting, relocating the equipment or shielding the location.

For more information on how to manage the unit's RF electromagnetic environment, please consult the AMI TIR 18-1997 - Guidance on Electromagnetic Compatibility of Medical Devices for Clinical/Biomedical Engineers.

Electromagnetic Emissions

Guidance and Manufacturer's Declaration - Electromagnetic Emissions - For all Equipment and Systems

The *Voyager* is intended for use in the electromagnetic environment indicated below. The customer or the user of the *Voyager* should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment - guidance	
RF emissions CISPR 11	Group 1	The <i>Voyager</i> uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.	
RF emissions CISPR 11	Class B	The <i>Voyager</i> is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.	
Harmonic emissions	Class A		
IEC 61000-3-2			
Voltage fluctuations/flicker emissions IEC 61000-3-3	Complies		

Electromagnetic Compatibility

Electromagnetic Immunity

Guidance and Manufacturer's Declaration - Electromagnetic Immunity - For all Equipment and Systems

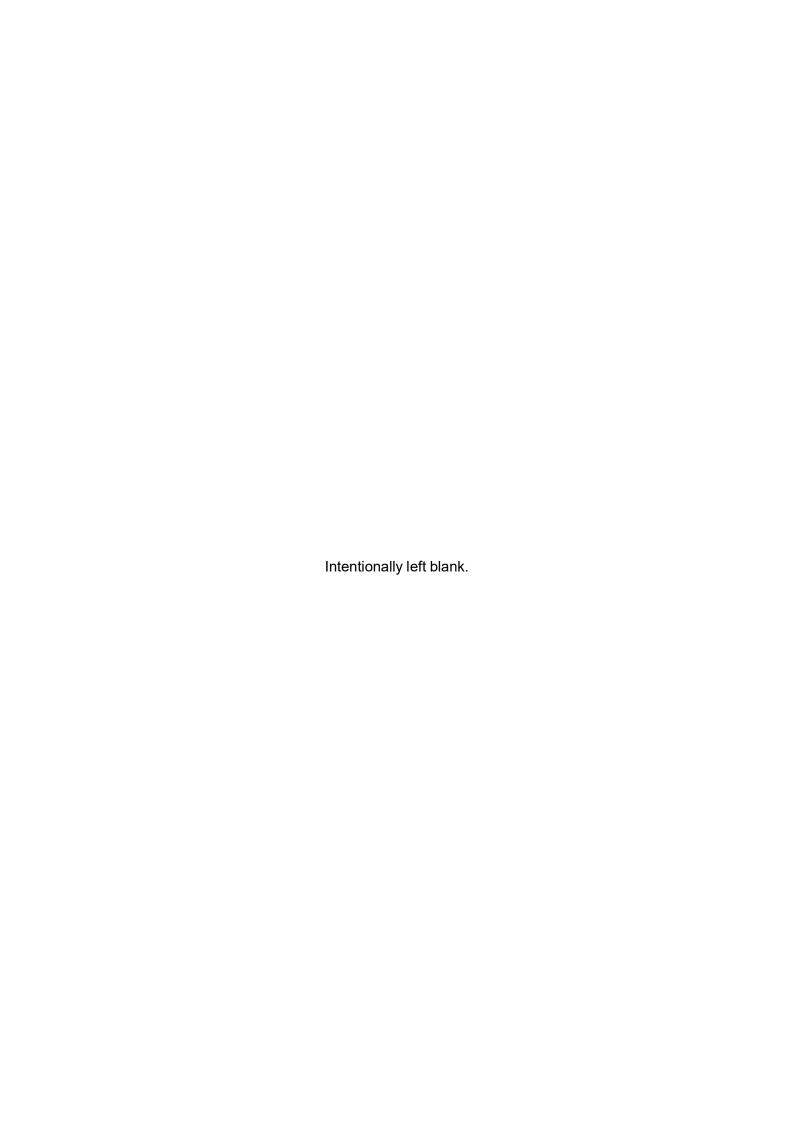
The *Voyager* is intended for use in electromagnetic environment specified below. The customer or the user of the *Voyager* should assure that it is used in such an environment.

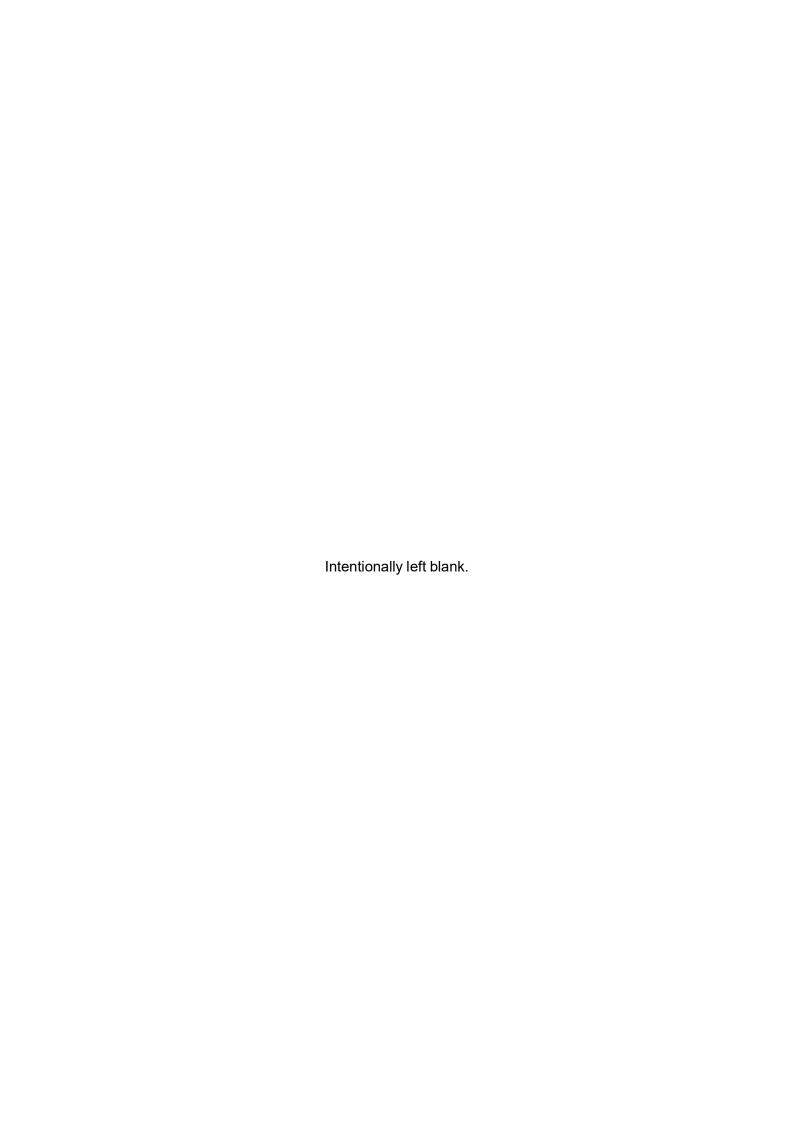
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - Guidance	
Electrostatic discharge (ESD)	±8 kV contact ±15 kV air	±8 kV contact ±15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.	
Electrical fast transient/burst	±2 kV, AC Mains	±2 kV, AC Mains	Mains power quality should be that of a typical commercial or hospital environment.	
	±1 kV, I/O Ports	±1 kV, I/O Ports		
IEC 61000-4-4	100 kHz repetition frequency	100 kHz repetition frequency		
Surge	±2 kV, AC Mains, Line to Ground	±2 kV, AC Mains, Line to Ground	Mains power quality should be that of a typical commercial or hospital environment.	
IEC 61000-4-5	±1 kV, AC Mains, Line to Line	±1 kV, AC Mains, Line to Line		
Voltage dips, short interruptions and voltage variations on power supply input lines	0 % <i>U</i> τ; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0 % <i>U</i> τ; 1 cycle and 70 % <i>U</i> τ; 25/30 cycles Single phase: at 0° 0 % <i>U</i> τ; 250/300 cycle	0 % <i>U</i> T; 0,5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0 % <i>U</i> T; 1 cycle and 70 % <i>U</i> T; 25/30 cycles Single phase: at 0° 0 % <i>U</i> T; 250/300 cycle	Mains power quality should be that of a typical commercial or hospital environment.	
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m 50/60 Hz C mains voltage prior to appl	30 A/m 50/60 Hz	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercials or hospital environment.	

Electromagnetic Compatibility

(continued)

Guidance and Manufacturer's Declaration - Electromagnetic Immunity - For all Equipment and Systems						
Immunity test	IEC 60601 test level	Compliance level	Electromagnetic environment - guidance			
Conducted RF IEC 61000-4-6	3 V outside ISM bands between 0.15-80 MHz 6 V inside ISM and	3 V outside ISM bands between 0.15-80 MHz 6 V inside ISM and amateur radio bands	N/A			
	amateur radio bands between 0.15-80 MHz	between 0.15-80 MHz				
Radiated RF IEC 61000-4-3	10 V/m 80 MHz to 2.7 GHz	10 V/m 80 MHz to 2.7 GHz	N/A			
Proximity fields from RF wireless communications equipment IEC 61000-4-3	380 - 390 MHz 27 V/m; PM 50 %; 18 Hz 430 - 470 MHz 28 V/m; (FM ±5 kHz, 1 kHz sine) PM; 18 Hz 800 - 960 MHz 28 V/m; PM 50 %; 18 Hz 1700 - 1990 MHz 28 V/m; PM 50 %; 217 Hz 2400 - 2570 MHz 28 V/m; PM 50 %; 217 Hz 5100 - 5800 MHz 9 V/m; PM 50 %; 217 Hz	380 - 390 MHz 27 V/m; PM 50 %; 18 Hz 430 - 470 MHz 28 V/m; (FM ±5 kHz, 1 kHz sine) PM; 18 Hz 800 - 960 MHz 28 V/m; PM 50 %; 18 Hz 1700 - 1990 MHz 28 V/m; PM 50 %; 217 Hz 2400 - 2570 MHz 28 V/m; PM 50 %; 217 Hz 5100 - 5800 MHz 9 V/m; PM 50 %; 217 Hz	N/A			
	704 - 787 MHz 9 V/m; PM 50 %; 217 Hz	704 - 787 MH32z 9 V/m; PM 50 %; 217 Hz				





AUSTRALIA Arjo Australia Building B, Level 3 11 Talavera Road

Macquarie Park, NSW, 2113,

Australia

Phone: 1800 072 040

BELGIQUE / BELGIË Arjo Belgium Evenbroekveld 16 9420 Erpe-Mere

Phone: +32 (0) 53 60 73 80 Fax: +32 (0) 53 60 73 81 E-mail: info.belgium@arjo.com

BRASIL

Arjo Brasil Equipamentos Médicos Ltda Rua Marina Ciufuli Zanfelice, 329 PB02

Galpão - Lapa

São Paulo - SP - Brasil CEP: 05040-000 Phone: 55-11-3588-5088

E-mail: vendas.latam@arjo.com E-mail: servicios.latam@arjo.com

CANADA

Arjo Canada Inc.

90 Matheson Boulevard West

Suite 300

CA-MISSISSAUGA, ON, L5R 3R3

Tel/Tél: +1 905 238 7880

Free: +1 800 665 4831 Institutional Free: +1 800 868 0441 Home Care

Fax: +1 905 238 7881 E-mail: info.canada@arjo.com

ČESKÁ REPUBLIKA Arjo Czech Republic s.r.o.

Na Strzi 1702/65 140 00 Praha Czech Republic

Phone No: +420225092307 e-mail: info.cz@arjo.com

DANMARK

Arjo A/S

Vassingerødvej 52 DK-3540 LYNGE Tel: +45 49 13 84 86 Fax: +45 49 13 84 87

E-mail:

dk_kundeservice@arjo.com

DEUTSCHLAND

Arjo GmbH

Peter-Sander-Strasse 10 DE-55252 MAINZ-KASTEL Tel: +49 (0) 6134 186 0 Fax: +49 (0) 6134 186 160 E-mail: info-de@arjo.com

ESPAÑA

ARJO IBERIA S.L.
Poligono Can Salvatella
c/ Cabanyes 1-7
08210 Barberà del Valles
Barcelona - Spain

Telefono 1: +34 900 921 850 Telefono 2: +34 931 315 999 FRANCE Arjo SAS

2 Avenue Alcide de Gasperi

CS 70133

FR-59436 RONCQ CEDEX Tél: +33 (0) 3 20 28 13 13 Fax: +33 (0) 3 20 28 13 14 E-mail: info.france@arjo.com

HONG KONG

Arjo Hong Kong Limited

Room 411-414, 4/F, Manhattan Centre, 8 Kwai Cheong Road, Kwai Chung, N.T.,

HONG KONG Tel: +852 2960 7600 Fax: +852 2960 1711

ITALIA

Arjo Italia S.p.A. Via Giacomo Peroni 400-402 IT-00131 ROMA

Tel: +39 (0) 6 87426211 Fax: +39 (0) 6 87426222 E-mail: Italy.promo@arjo.com

MIDDLE EAST

Arjo Middle East FZ-LLC
Office 908, 9th Floor,
HQ Building,North Tower,
Dubai Science Park,
Al Barsha South
P.O Box 11488, Dubai,
United Arab Emirates
Direct +971 487 48053
Fax +971 487 48072

Email: Info.ME@arjo.com

NEDERLAND

Arjo Nederland BV Biezenwei 21 4004 MB TIEL Postbus 6116 4000 HC TIEL

Tel: +31 (0) 344 64 08 00 Fax: +31 (0) 344 64 08 85 E-mail: info.nl@arjo.com

NEW ZEALAND

Arjo Ltd 34 Vestey Drive Mount Wellington NZ-AUCKLAND 1060 Tel: +64 (0) 9 573 5344 Free Call: 0800 000 151 Fax: +64 (0) 9 573 5384

E-mail: nz.info@Arjo.com

NORGE

Arjo Norway AS Olaf Helsets vei 5 N-0694 OSLO Tel: +47 22 08 00 50

Faks: +47 22 08 00 51

E-mail: no.kundeservice@arjo.com

ÖSTERREICH

Arjo GmbH

Lemböckgasse 49 / Stiege A / 4.OG

A-1230 Wien Tel: +43 1 8 66 56 Fax: +43 1 866 56 7000 **POLSKA**

Arjo Polska Sp. z o.o. ul. Ks Piotra Wawrzyniaka 2 PL-62-052 KOMORNIKI (Poznań)

Tel: +48 61 662 15 50 Fax: +48 61 662 15 90 E-mail: arjo@arjo.com

PORTUGAL

Arjo em Portugal MAQUET Portugal, Lda. (Distribudor Exclusivo) Rua Poeta Bocage n.º 2 - 2G PT-1600-233 Lisboa Tel: +351 214 189 815 Fax: +351 214 177 413 E-mail: Portugal@arjo.com

SUISSE / SCHWEIZ

Arjo AG
Fabrikstrasse 8
Postfach

CH-4614 HÄGENDORF Tél/Tel: +41 (0) 61 337 97 77 Fax: +41 (0) 61 311 97 42

SUOMI

Arjo Scandinavia AB Riihitontuntie 7 C 02200 Espoo Finland

Puh: +358 9 6824 1260

E-mail: Asiakaspalvelu.finland@arjo.com

SVERIGE

Arjo International HQ Hans Michelsensgatan 10 SE-211 20 MALMÖ Tel: +46 (0) 10 494 7760 Fax: +46 (0) 10 494 7761 E-mail: kundservice@arjo.com

UNITED KINGDOM

Arjo UK and Ireland Houghton Hall Park Houghton Regis UK-DUNSTABLE LU5 5XF

Tel: +44 (0) 1582 745 700 Fax: +44 (0) 1582 745 745 E-mail: sales.admin@arjo.com

USA

Arjo Inc.

2349 W Lake Street Suite 250 US-Addison, IL 60101 Tel: +1 630 307 2756

Free: +1 80 0 323 1245 Institutional Free: +1 800 868 0441 Home Care

Fax: +1 630 307 6195 E-mail: us.info@arjo.com

JAPAN

Arjo Japan K.K. 東京都港区虎ノ門三丁目7番8号 ランディック第2虎ノ門ビル9階

Tel: +81 (0)3-6435-6401 Fax: +81 (0)3-6435-6402 E-mail: info.japan@arjo.com

At Arjo, we are committed to improving the everyday lives of people affected by reduced mobility and age-related health challenges. With products and solutions that ensure ergonomic patient handling, personal hygiene, disinfection, diagnostics, and the effective prevention of pressure ulcers and venous thromboembolism, we help professionals across care environments to continually raise the standard of safe and dignified care. Everything we do, we do with people in mind.



ArjoHuntleigh AB Hans Michelsensgatan 10 211 20 Malmö, Sweden www.arjo.com





