CHANGING PLACES CARE ENVIRONMENTS HYGIENE ROOMS





Instruction Manual



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Introduction

Application

The primary purpose of a ceiling lift system is to safely lift and transfer a patient with as little effort as possible for the caregiver, regardless of the room type. A ceiling lift is simple and safe for the caregiver as well as the patient. The lift systems fit into all environments.

Description

The GoLift is an ideal ceiling lift system designed for routine transfers of patients. The most compact ceiling lift in its class, the GoLift is designed to be aesthetically pleasing to both the caregiver and the patients. In an effort to address infection control requirements, we gave careful consideration to the smooth edges and rounded corners of our ceiling lift, carry bar and hand control. When you look under the cover of this compact lift, you will find an impressive set of all metal gears and state-of-the-art battery technology that will allow the caregiver to safely, and effortlessly, transfer a patient weighing up to 700 lbs on a single lift. For larger patients, you may combine two compact GoLift systems to obtain a lifting weight of 1000 lbs.. In addition, our revolutionary trolley design allows for quick installation of the lift into and out of the track. The trolley provides an instant mechanical and electrical connection while our modular track system gives you tremendous flexibility so that your workspace may be optimised to suit your needs for any working environment.

The GoLift is available in three weight capacities which must not be exceeded:

400 lbs (182 kg) 450 lbs (205 kg) 700 lbs (318 kg)

The GoLift is a fixed lift and can be installed in a variety of track profiles (the trolley can be customised to fit these track profiles)

Contents of Packaging:

- 1. GoLift
- 2. Hand Control
- 3. Charger
- 4. Owner's Manual
- 5. GoLift Trolley
- NOTE: The Carry Bar is packaged separately.

Upon receipt of the packages, verify it against the packing slip to ensure the shipment is complete and inspect the equipment for possible damage. If there is any damage, DO NOT USE the equipment and notify the carrier immediately to file a claim. Provide complete information concerning damage claims or shipping errors to Wealden Rehab. Include all equipment identification numbers along with a description of the damaged parts.





Symbols used

Symbol	Reference	Title
	ISO 7000-0434A	Caution risk of danger
C US	TUV	Certified by TUV
📩 Туре В	IEC 60417-5840	Type B Applied Part
CE	CE	Certification of Conformity
I	ISO 7010-M002	Refer to instruction manual/booklet
	AM	Emergency Lowering
	N/A	N/A



WARNING: This symbol is intended to alert the user of hazard or unsafe practices, which could result in serious bodily harm.

Markings

The GoLift 400 and GoLift 700 are designed to comply with the following Standards:

Models:	400 lbs and 700 lbs
Brand Name:	GoLift
Product:	GoLift
Standard(s):	CAN/CSA-C22.2 No. 60601-1:08 Medical Electrical Equipment – Part 1: General requirements for basic safety and essential performance. ISO10535:2006 Hoists for the transfer of disables persons – Requirements and Test Methods





Safety Instructions



Read these instructions carefully or serious injury may occur

- The GoLift must be installed only by authorised personnel
- Do not use this equipment prior to understanding the contents of this manual.
- Contents of this manual are subject to change without prior notice to users. Keep for future reference.
- Never place the GoLift, track/PLP and sling(s) in control of a person who has not been properly trained in the use and care of this equipment.
- The GoLift and associated Track/PLP and sling(s) are for transferring patients only. Never use the GoLift for any other purpose.
- Wealden Rehab's Warranty is void if unauthorised personnel perform service on the GoLift system.
- In facilities where more than one caregiver is be responsible for using the GoLift and associated track and slings, it is important that all caregivers be trained in the proper use of this equipment. A training program should be established by the facility to familiarise new caregivers with this equipment.
- Do not expose the GoLift directly to water. Warranty does not cover any misuse or abuse of the GoLift.
- The GoLift should be inspected and maintained on a regular basis to keep it operating safely and correctly. Refer to page 16 of this manual.
- Any accessories used with the GoLift including the track/PLP and sling(s) should be checked to ensure that they are in good working order. Check for signs of wear or fraying prior to use. Report any unusual wear or damage immediately.
- Wealden Rehab will not be responsible for any damage caused by misuse, neglect or purposeful destruction of the lift and its associated components. Do not attempt to modify/alter the GoLift.
- Do not in any circumstance exceed the maximum allowable load of this lift. Refer to the "Technical Specifications" section of this manual and/or the labels on the lift.
- The installation of the lift, track and sling are certified to a maximum load. Do not exceed the maximum rated load of any of the components.
- There is a risk of explosion if the lift is used in the presence of flammable anesthetics.
- The GoLift should be decommissioned/disposed of after the recommended service lift in accordance with local law regulations.



There are no known contraindications associated with the use of the Amico GoLift and its accessories, provided they are used per our recommendations and guidelines.

However, for any independent uses of the GoLift, it is extremely important that the patient is able to receive assistance, during the transfer in the event of an equipment failure. This assistance can be provided in the form of; a nearby qualified caregiver, a phone or other communication device.



Technical Specifications for GoLift

Four GoLift Weight Capacities





Safe Working Load (SWL)

400lbs / 181kg 450lbs / 205kg 700lbs / 318kg 1000lbs / 454kg

Dimensions

400, 450 & 700lbs / 181, 205 & 318kg:

- Length: 7 ³/₄" / 197 mm
- Width: 7 3/4" / 197 mm
- Height: 4 ¼" / 108 mm
- Strap Length: 84" / 2134 mm

1000 lbs / 454 kg:

- Length: 15 1/2" / 394 mm
- Width: 7 3/4" / 197 mm
- Height: 4 ¼" / 108 mm
- Strap Length: 84" / 2134 mm

Unit Weight

400lb / 181kg version: 8lbs / 4kg 450lb / 205kg version: 10lbs / 5kg 700lb / 318kg version: 10lbs / 5kg 1000lb / 454kg version: 20lbs / 9kg (Tandem Lifts)

Lift Case Flame retardant ABS

Safety

Emergency Stop Emergency Lowering Device Upper Limit Detection Lower Limit Detection Slack Tape Sensor Free Fall Brake (over speed governor) Low Battery and Dead Battery Alarms Soft Start and Stop Overload Protection Emergency Manual Lowering

Approvals

Certified to: Can/CSA-C22.2 No 60601-1:08, UL 60601-1:08 Tested to: ISO 10535-06, CE

Hand Control

Capacitive Touch Protection Class: IPX0

Service Life

10 years or 22,500 cycles

Maximum Sound Level <65 dB

Maximum Lifting Speed

No load: 2 inches/second 150lbs / 68kg: 1.3 inches/second 400lb / 181kg: 1 inch/second 700lb / 318kg: 1 inch/second 1000lb / 454kg: 1 inch/second

Batteries

High Capacity, Nickel Metal Hydride (Ni-Mh) Standard: 2 x 14.4V (2Ah) Optional: 2 x 14.4V (3.3Ah) or (5Ah)

Battery Charger:

100-240 VAC 40 Watt

Number of Lifts per Charge (Duty: 10/90)

- 25% of strap at midrange¹:
 - 375 with 185lbs / 84kg
 - 170 with 400lbs / 181kg
 - 102 with 700lbs / 318kg

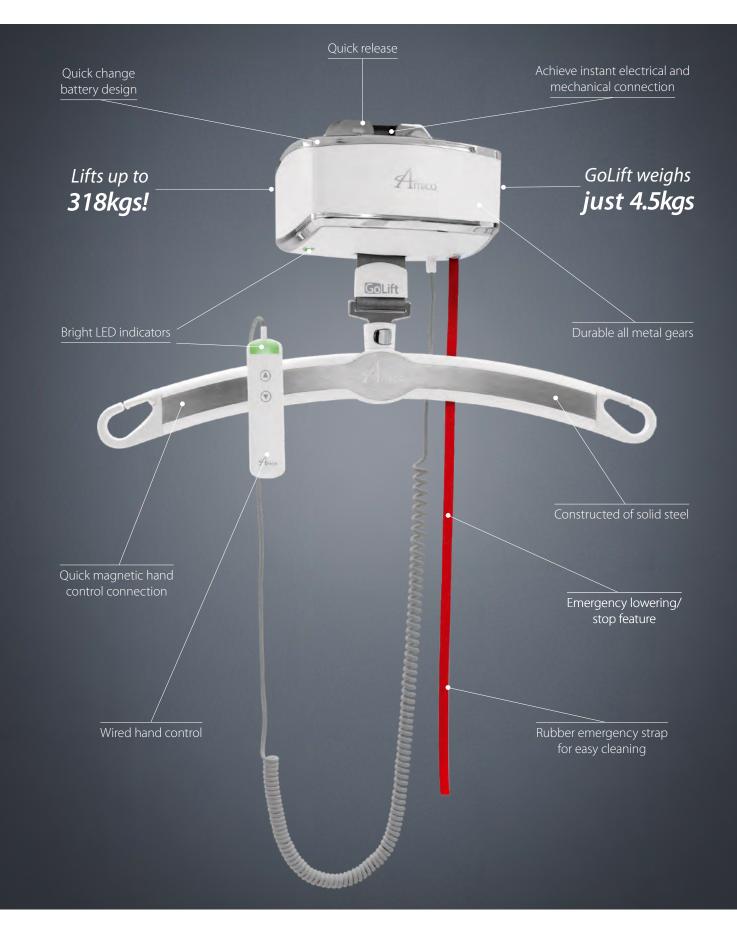
Charging time: 2 - 4 hours

¹Calculated using 5Ah Packs



GOLift - INSTRUCTION MANUAL

Anatomy of the GoLift





Basics in Transferring a Patient

Lifting Sling

A lifting sling with four to six straps designed for mounting on hooks should be used when using a GoLift Carry Bar. Place the straps on the hooks and make sure the loops are not in the latch so the straps do not

unintentionally fall off.

Wealden Rehab shall not be liable for faults or accidents due to incorrect use of the lifting sling, or for reasons of inadequate attention on the part of the caregiver or patient.

Working with the GoLift

The GoLift moves freely in the track system and does not have any special requirements for space or power in connection with moving. Attention can this be fully focused on the user's functional level and the caregivers technique.

To use the GoLift correctly, the patient should only be lifted to the extent that she/he is clear of the surface and should be moved at this height.

Attaching the Lifting Sling

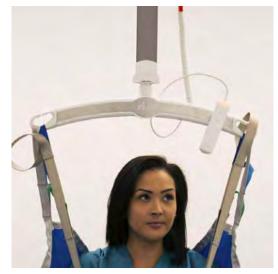
Place the straps from the lifting sling on the hooks on the carry bar. Start with the uppermost set of straps (from the back) and then take the lowest set of straps (from the legs)



Be careful when attaching the lifting sling on the hooks. Check that the straps have been completely through the opening and into place in the carry bar hooks. When pressing the up button to lift the patient, check again that all straps remain correctly placed in the carry bar's hooks.

Lifting to and from a seated position

- When lifting a patient from e.g. a wheelchair, move the GoLift towards the patient to be lifted.
- The carry bar should be at the same height as the patient's chest and should not be moved further in over the user to approximately mid-though position.
- Place the carry bar parallel to the patient's shoulders.
- Place the lifting sling behind the user's back between the back of the chair and the user's back.
- The center band of the lifting sling should follow the user's spine. Lead the leg straps along the outer sides of the patient's shins and beneath the thighs between the hollow of the knees and the hip joints. Cross the leg straps in front of the user.
- All four lifting straps are now ready to be attached. The lifting sling can now be mounted on the carry bar.







Basics in Transferring a Patient

Lifting to and from lying position in bed:

- Bring the carry bar over the center of the patient to be lifted.
- Place the carry bar parallel to the patient's shoulders.
- Turn the patient onto his or her side. The sling should be placed so that the top of the sling is at the same height as the top of the user's head. Now position the sling over the user so that the center band follows the user's spine. Turn the user onto his or her back and pull out the remaining part of the lifting sling. Place the leg straps beneath the user's thighs and cross them. All four lifting straps are now ready to be attached and the lifting sling can now be mounted on the carry bar. It is an advantage to elevate the head of the bed so that the patient is sitting up.
- Only persons who have received competent instruction regarding the use of the lifting equipment and fitting of slings should use the GoLift.



Important: Plan the move and avoid leaving the patient in the sling unattended. Before lifting, check that the patient is completely free of his/her surroundings. The patient's head, arms, hands and feet must not be in danger or becoming trapped. Be careful with any tubes and wires that are attached to the user. Check that the hand control and hand control cable is free of hanger, patient and other object before the lift is activated up or down moved.

Installing the GoLift in PLP Track

- 1. Remove both the end caps on the PLP Arm.
- Make sure the trolley is inserted correctly in the GoLift.
 The trolley is equipped with sensors that will only allow the GoLift to operate when the trolley is secured inside the lift.









Installing the GoLift in PLP Track

3. By using an adjustable wrench, remove the end stopper on the PLP track by loosening the bolts.

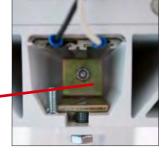
4. Slide the GoLift motor into the PLP track from the other side. Make sure the GoLift Motor is constantly in contact with the charging strip.

Charging Strip

- 5. Fasten the end stops back on to the PLP track. Ensure there is enough clearance room to place the end cap on the PLP arm and place the end cap back on the arm.
- 6. Fasten all end stops tightly using an adjustable wrench.
 - NOTE: If you are installing the GoLift in an existing track system you must ensure that the max load of the track system is equal or higher than the max load of the GoLift.
- 7. To disconnect the lift from the trolley, press on the two quick release trolley buttons and pull down on the lift.













Installing the Endstop on the GoLift Track

1. Use a 3/16 Allen key to secure the endstop to the track.

GoLift Track Endstop



Connecting the Carry Bar to the Lift Strap

- 1. Hold the carry bar and press the button using the thumb as shown.
- 2. Insert the strap attachment in the slot on the carry bar top cover with the open side facing down and release the button.
- 3. Check that the button has returned to its locked position by checking that it is flush with the cover of the carry bar and that the strap attachment can rotate freely.







Connecting the Hand Control to the Carry Bar or Wall Plate

Capacitive touch hand control – Allows users with limited dexterity to effortlessly operate the unit.

Magnetic Hand control connect for quick attachment to carry bar or wall plate.



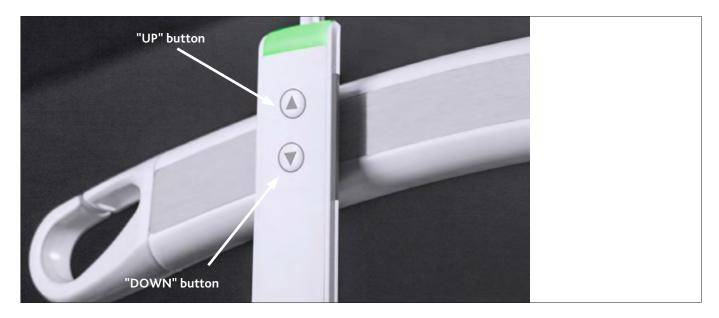
Do not place the hand control within 5" of a pacemaker. Patients with pacemakers must follow the instructions provided by their doctors.





Operating the GoLift

The GoLift is switched on automatically when a button on the hand control is pressed. The GoLift switches off automatically after approximately three minutes without activation.



Charging the GoLift

The charger contacts with two metal charging strips located inside the track.

Whenever the lift is over a section of track with charging strips, it will automatically start to charge the lift if the battery is low.

The batteries should be charged on a regular basis. It is recommended that the lift be left on charge when not in operation, and at the end of each day. This will maximise the life cycle of the batteries.

The GoLift may remain connected to the charger indefinitely since the charger has a builtin regulator, eliminating the danger of overcharging.

As a general rule, it is recommended that the carry bar be raised to a height that will not interfere with anything or anyone while the lift is not in use.





Do not drain the batteries excessively. This will dramatically reduce the lifespan of the batteries. If the slow beep sounds, be sure to recharge the battery as soon as possible.



Do not leave the GoLift with the power on for a long time. If the GoLift is not returned to the charger the batteries will be drained and damaged.



Do not install the power supply in a position where it is not possible to disconnect the plug from the charger.

NOTE: Ensure that the trolley is in the charging section of the track when installing the lift.



LED and Buzzer Functions

Charging	Battery	LED	Buzzer	Additional Notes	
	Green				
Yes	Full	If re-connect to charger LED goes Orange for 5 seconds then back to Green	N/A	LED stays on 1 minute after use then shuts off	
No	Full	Green	N/A	LED stays on 1 min then shuts off	
Yes	Not Full	Orange	N/A	LED stays on 1 min then shuts off	
No	Low Battery	Blinks Orange	N/A	LED stays on 1 min after use then shuts off. Monitor Voltage to check for battery level. 1 full lift at max load must be able to be completed.	
No	Dead	Red	Single beep every 3 seconds	LED stays on 1 min after use then shuts off. Monitor Voltage to check for battery level. Lift in up direction not allowed but must be able to go in the down direction.	
	Fault		Beep pattern		
No	Temperature Error	Flashing Red	single beep every 5 seconds	LED stays on for 1 minute	
Emergency Lowering Red		Веер	Red LED if main board is working. No LED if main board has malfunctioned.		
Latch ErrorRed flashing once every .5 secondBuzzer On solid					
During Sleep mode (when LED turns off - after 1 minute) the unit should draw no power (1ma).					
0				are pressed the unit has to wake up instantly.	

Maintenance Alarm

At 1001 lifts, the LED will flash Green.

To reset, press Up button, then Down button, then Up button, then Down button and then hold both buttons for 10 seconds.

If successful, the LED turns Red, Green and Orange. The buzzer beeps three times.

Please note, that a reset can be performed at any time, not just after the 1000th lift.

Emergency Stop

The GoLift unit also has an Emergency shut-off feature that allows the operator to shut the power to the lift completely in the event of an emergency. By pulling once on the RED emergency lowering cord, located on the underside of the lift, the lift will immediately stop and all its functions will be disabled. The "ON" indicator light will turn off, and the Emergency Shut-off button located inside the lift will pop out. After an emergency, the lift must be inspected prior to restoring to use. In order to restore power to the lift, the tab must be pressed back into the lift.





Do not pull the red cord forcefully. For assistance after an emergency contact: Wealden Rehab on 01634 813388



Emergency Lowering

In the event that the DOWN button on the hand control does not function, or in power failure situations, the patient may be lowered by pulling and holding down the RED emergency lowering cord located on the underside of the lift.

Continue to pull down until the patient is safely lowered to the desired position. The lift will beep as you continue to pull down on the cord and will continue beeping until the cord is released after the desired lowering has beenachieved.

NOTE: The emergency lowering button does not provide a raising function. The failure of any of the lowering

devices should be reported to Wealden Rehab.



In an event of emergency when normal lowering system of the lift malfunctions and the "emergency lowering" function is used, the lift must be reset by a qualified lift technician before re-use.

Manual Emergency Lowering

The manual emergency and raising should only be used if the Emergency Lowering does not work. A proper safety ladder of stool may be required to remove the plug from the cover.



Caution: DO NOT attempt to use the lift while using manual lowering.

Remove the round plug from the lift cover and use 3/16" or 2.5 mm Allen key to rotate the motor in the up or down direction.



Overspeed Cam

The Overspeed Cam brake is made of a metal bar fixed to the drum. Incase of gear or motor breakage, the centrifugal force created will block the bar against the frame.



Cleaning and Disinfection

The exterior of the GoLift should only be cleaned, disinfected using the recommended cleaning agents shown below. Damp a cloth with the cleaning agent and wipe down entire exterior of the lift and carry bar. Other chemicals and/or liquids not listed should not be used to clean and disinfect this lift.



Take great care to ensure that no liquids get inside the GoLift. The lift is not drip proof or water tight. Failure to protect the lift from liquids may result in damage to the lift and may cause personal injury.

Recommended cleaning agents:

- Virox Accel TB
- Virox 5
- Dispatch Hospital Cleaner disinfectant towels with Bleach
- Clorox Healthcare Professional Disinfecting Bleach Wipes
- Sani-Cloth super germicidal disposable wipes
- Virocidin-X

Troubleshooting

Should problems arise with the use of the GoLift, review the following chart. Find the fault and complete the recommended solution. If the fault is not found and or/the solution does not correct the problem contact Wealden Rehab.

Fault	Recommended Solution
The GoLift emits a slow beep while in use. The lift only goes down but not in the upwards direction.	The batteries are low and the lift should be charged.
The GoLift cannot lift.	If the load is in excess of the safe working load the GoLift will not work due to over-current protection.
The batteries are always dead after a few charges	Replace the batteries as they may be at the end of their service life
The GoLift does not operate when you press the buttons on the hand control.	Check that the emergency cord is not pulled.
The GoLift is not operating properly	Make sure the GoLift is inserted all the way inside the trolley. This would activate the trolley's limit switches.
The lift does not go up	There may be a twist in the lift strap.



Inspection and Maintenance

Prior to using the GoLift, the inspections should be conducted per the following schedule:

ltem	Before Use	Every Month	BiAnnual	Every Two Years
GoLift	•			
Ensure that end stops are installed	•			
Inspect strap for wear or fraying	•			
Ensure Batteries are charged	•			
Inspect carry bar for damage or sharp edges			•	
Inspect the wheels in the trolley. Replace if damaged.			•	
Sling				
Check all straps for wear or fraying	•			
Inspect the sling for any damage in the fabric	•			
Ensure there are no loose threads in the stitching	•			
Maintenance by a certified technician Replace the strap			•	•
Inspect the gearbox and carry bar (check for cracks)			•	
Inspect the gears for any broken or worn teeth			٠	
** Verify the overspeed cam is operating freely			•	
Check emergency stop cord			•	
** Check emergency lowering device			٠	
*Annual load test with SWL (maximum safe working load)			٠	
Ensure the end stops are installed			•	

- * In accordance to the ISO 10535 Standard "Hoists for the transfer of disabled persons Requirements and test methods" an inspection should be performed on the GoLift at least once a year. This inspection should be performed by a qualified technician and should include a working load test of one (1) lifting cycle with the maximum load.
- ** These two functions must be checked by a qualified technical to ensure the essential performance of the GoLift.



Do not operate the GoLift until any issues discovered during the inspection have been addressed by a certified technician.



GoTrack Systems and Support Structure

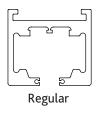
GoTrack Configurations

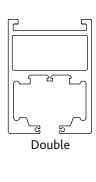
We offer three different track profiles, as well as modular and custom configurations that come in:

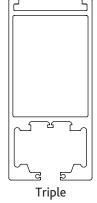
- Straight Track
- Curved Systems
- Full Room Coverage (X/Y or H Systems)
- Optional Infection Control Track
- Optional recessed, flush mount track systems

All GoTrack Systems feature:

- Low weight and high capacity
- Fast and flexible installation
- Hidden End Stops for safety
- Anywhere charge







Description	Part Number	Length x Width x Height	Weight per metre	Material	
Regular	TRK-REG-120	120 x 2.63 x 2.36" / 3000 x 67 x 60 mm	8.3 lbs / 3.77 kg	White powder coated extruded aluminum	
Regular	TRK-REG-150	150 x 2.63 x 2.36" / 3800 x 67 x 60 mm	8.3 lbs / 3.77 kg	White powder coated extruded aluminum	
Regular	TRK-REG-180	180 x 2.63 x 2.36" / 4500 x 67 x 60 mm	8.3 lbs / 3.77 kg	White powder coated extruded aluminum	
Regular	TRK-REG-240	240 x 2.63 x 2.36" / 6100 x 67 x 60 mm	8.3 lbs / 3.77 kg	White powder coated extruded aluminum	
Double	TRK-DBL-120	120 x 2.63 x 3.55" / 3050 x 67 x 90 mm	10.4 lbs / 4.72 kg	White powder coated extruded aluminum	
Double	TRK-DBL-150	150 x 2.63 x 3.55" / 3800 x 67 x 90 mm	10.4 lbs / 4.72 kg	White powder coated extruded aluminum	
Double	TRK-DBL-180	180 x 2.63 x 3.55" / 4500 x 67 x 90 mm	10.4 lbs / 4.72 kg	White powder coated extruded aluminum	
Triple	TRK-TRP-156	156 x 2.63 x 5.75" / 4000 x 67 x 146 mm	12.4 lbs / 5.65 kg	White powder coated extruded aluminum	
Triple	TRK-TRP-240	240 x 2.63 x 5.75" / 6100 x 67 x 146 mm	12.4 lbs / 5.65 kg	White powder coated extruded aluminum	
Triple	TRK-TRP-315	315 x 2.63 x 5.75" / 8000 x 67 x 146 mm	12.4 lbs / 5.65 kg	White powder coated extruded aluminum	
Custom	Custom Custom lengths in any track type available upon request				



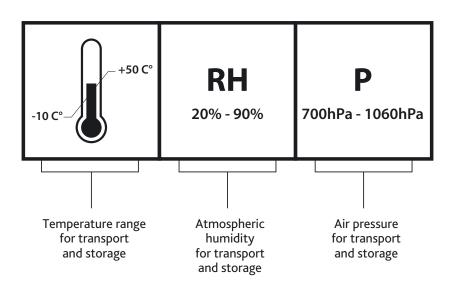
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Environmental Conditions

Operation				
Minimum Maximum				
Temperature	+10°C	+40°C		
Relative atmospheric humidity	30%	75%		
Air pressure	700 hPa	1060 hPa		

Transport/Storage				
Minimum Maximum				
Temperature	-10°C	+50°C		
Relative atmospheric humidity	20%	90%		
Air pressure	700 hPa	1060 hPa		

References on the Package





Guidance and Manufacturer's Declaration - Electromagnetic Emissions

The [EQUIPMENT or SYSTEM] is intended for use in the electromagnetic environment specified below. The customer or the user of the [EQUIPMENT or SYSTEM] should assure that it is used in such an environment.

Emissions Test	Compliance	Electromagnetic environment – guidance
RF emissions CISPR 11	Group 1	The [EQUIPMENT or SYSTEM] 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 A	The [EQUIPMENT or SYSTEM] is suitable for use in all establishments other than domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations / flicker emissions IEC 61000-3-3	Complies	



Guidance and Manufacturer's Declaration - Electromagnetic Emissions

The [EQUIPMENT or SYSTEM] is intended for use in the electromagnetic environment specified below. The customer or the user of the [EQUIPMENT or SYSTEM] should assure that it is used in such an environment.

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance
Electrostatic discharge (ESD) IEC 61000-4-2	±6 kV contact	±6 kV contact	Floors should be wood, concrete or ceramic tile. If floors are covered
	±8 kV air	±8 kV air	with synthetic material, the relative humidity should be at least 30 %
Electrical fast transient/burst	±2 kV for power supply lines	±1 kV for power supply lines	Mains power quality should be that
IEC 61000-4-4	±1 kV for input/ output lines	±0.250 kV for input/output lines	of a typical commercial or hospital environment.
Surge IEC 61000-4-5	±1 kV line(s) to line(s)	±1 kV line(s) to line(s)	Mains power quality should be tha
	±2 kV line(s) to earth	±2 kV line(s) to earth	of a typical commercial or hospita environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<5 % UT (>95 % dip in UT) for 0,5 cycle 40 % UT (60 % dip in UT) for 5 cycles 70 % UT (30 % dip in UT) for 25 cycles <5 % UT (>95 % dip in UT) for 5 sec	<5 % UT (>95 % dip in UT) for 0,5 cycle 40 % UT (60 % dip in UT) for 5 cycles 70 % UT (30 % dip in UT) for 25 cycles <5 % UT (>95 % dip in UT) for 5 sec	Mains power quality should be that of a typical commercial or hospital environment. If the user of the [EQUIPMENT or SYSTEM] requires continued operation during power mains interruptions, it is recommended that the [EQUIPMENT or SYSTEM] be powered from an interruptible power supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	3 A / m	Not Applicable	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.



Gu	Guidance and Manufacturer's Declaration - Electromagnetic Emissions				
The [EQUIPMENT or SYSTEM] is intended for use in the electromagnetic environment specified below. The customer or the user of the [EQUIPMENT or SYSTEM] should assure that it is used in such an environment.					
Emissions Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment – Guidance		
Field of the second se	3 Vrms 150 kHz to 80 MHz 3 V/m 80 MHz to 2,5 GHz	3 Vrms 3 V/m	Electromagnetic Environment – Guidance Portable and mobile RF communications equipment should be used no closer to any part of the [ME EQUIPMENT or ME SYSTEM], including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance $d = 1.2 \sqrt{P}$ $d = 1.2 \sqrt{P}$ 80 MHz to 800 MHz $d = 2.3 \sqrt{P}$ 800 MHz to 2,5 GHz where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation distance in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, a. should be less than the compliance level in each frequency range. b. Interference may occur in the vicinity of equipment marked with the following symbol:		

NOTE 1: At 80 MHz and 800 MHz, the higher frequency range applies.

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

a. Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the [ME EQUIPMENT or ME SYSTEM] is used exceeds the applicable RF compliance level above, the [ME EQUIPMENT or ME SYSTEM] should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the [ME EQUIPMENT or ME SYSTEM].

b. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.



Recommended separation distances between portable and mobile RF communications equipment and the [EQUIPMENT or SYSTEM]

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

Rated maximum output power of transmitter W	Separation distance according to frequency of transmitter M		
	150 kHz to 80 MHz d = 1.2√P	150 kHz to 80 MHz d = 1.2√P	800 MHz to 2,5 GHz d = 1.2√P
0,01	0.12	0.12	0.23
0,1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer. NOTE 1: At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies. NOTE 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

Disposal

- The GoLift doesn't contain any dangerous goods.
- The components of the GoLift should be properly disposed at the end of its shelf-life.
- Make sure that the materials are carefully separated.
- The electrical conducting boards should be submitted to an appropriate recycling facility.
- The rest of the components should be disposed according to the contained materials.



Warranty Policy - Mobility Solutions

The primary purpose of our ceiling lift system is to safely lift, transfer and reposition a patient with a little effort as possible for the caregiver, regardless of the room type. The Amico Mobility Solutions Corporation's GoLift is easy and safe to use for caregivers as well as patients. The lift systems fit into all environments and fulfills the highest requirements of function, safety and reliability.

Amico Mobility Solutions Corporation warrants its lifting equipment and workmanship to be free from defects for a period of three (3) years from the date of installation. This includes tracks, lift motor, and carry bar.

Within this period, Amico Mobility Solutions Corporation will replace any part (at no additional charge), which is deemed defective. Shipping and installation costs after the first twelve (12) months will be borne by the customer. The following exclusions apply: the warranty for batteries is for a period of three (3) months from the time of installation; the warranty for power supply is one (1) year from the time of installation.

This warranty is valid only when the product has been properly installed as outlined in the Amico Mobility Solutions Corporation specifications; including but not limited to proper usage and servicing of systems according to factory recommendations. It does not cover damages as a result of shipment failures, accidents, misuse, abuse, neglect, mishandling, alternation, misapplication or damages which may be attributed to acts of God.

Amico Mobility Solutions Corporation shall not be liable for incidental or consequential damages resulting from the use of the equipment.

All claims for warranty must first be approved by Amico Mobility Solutions Corporation's Service Department at acs-service@amico.com or 1.877.462.6426. A valid Return Goods Authorisation (RGA) number must be obtained from Amico Mobility Solutions Corporation prior to commencement of any service work. Warranty work which has not been per-authorised by Amico Mobility will not be reimbursed.





Speak to a **CareSpaces™** expert on 01634 949988, email carespaces@wealdenrehab.com or visit carespaces.co.uk

Trading as Wealden Rehab Ltd 113 Hopewell Drive, Chatham, Kent ME5 7NP United Kingdom











