



HOVERJACK® EMS Evacuation

User Manual



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HOVERJACK® EMS Evacuation User Manual

Symbol Reference



CE MARKING OF CONFORMITY



OPERATING INSTRUCTIONS



UK MARKING OF CONFORMITY



KEEP DRY



AUTHORIZED REPRESENTATIVE



LATEX FREE



UK RESPONSIBLE PERSON



MANUFACTURER



SWITZERLAND AUTHORIZED REPRESENTATIVE



DATE OF MANUFACTURE



CAUTION / WARNING



MEDICAL DEVICE



MANUAL CLEANING



MODEL NUMBER



IMPORTER



SERIAL NUMBER



DISPOSAL



DO NOT LAUNDER



FASTEN PATIENT SAFETY STRAP



UNIQUE DEVICE IDENTIFIER



FOOT END



PATIENT WEIGHT LIMIT



DECLARATION OF CONFORMITY

This product conforms to the requirements of Medical Devices Regulation (2017/745).

Intended Use and Precautions

INTENDED USE

The EMS Evacuation HoverJack® Device is used to lift a patient in supine position from the floor to bed or stretcher height, utilizing the HoverTech Air Supply to inflate each of the four chambers. The EMS Evacuation HoverJack can also be used to evacuate patients up or down stairs in the event of an emergency.

INDICATIONS

- Patients unable to assist in their own vertical lift, such as after a fall.
- Patients whose weight or girth poses a potential health risk for the caregivers responsible for lifting or evacuating said patients.
- CPR compatible when fully inflated.

CONTRAINDICATIONS

 Patients who are experiencing thoracic, cervical or lumbar fractures that are deemed unstable should not use the EMS Evacuation HoverJack unless a clinical decision has been made by your facility.

INTENDED CARE SETTINGS

 Hospitals, long term or extended care facilities, patient transport services and EMS.

PRECAUTIONS

- Make sure patient safety straps are secured before moving. Do not secure before inflation.
- Move the EMS Evacuation HoverJack using the transport straps and/or the transport handles located along the top perimeter.
- Never use patient safety straps to pull the EMS Evacuation HoverJack, as they may tear.
- When moving a patient on the inflated EMS Evacuation HoverJack, use caution and move slowly.

- Additional caregivers are recommended when moving or evacuating a patient over 350 lbs/159 kg.
- Never attempt to move a patient on an uninflated EMS Evacuation HoverJack.
- Never leave a patient unattended on an inflated device.
- Use this product only for its intended purpose as described in this
 manual
- Only use attachments and/or accessories that are authorized by HoverTech.

WARNING/CAUTION

- Always use a minimum of two caregivers while inflating/transporting the EMS Evacuation Hoverlack.
- A minimum of three caregivers is required for stairwell evacuation.
- One caregiver should remain on the side of the patient during inflation with the second caregiver managing inflation. If available, a third caregiver should attend to the other side of the patient.
- Reference product-specific user manuals for additional operating instructions.

PRECAUTIONS - HOVERTECH AIR SUPPLY

- Not for use in the presence of flammable anesthetics or in a hyperbaric chamber or oxygen tent.
- Route the power cord in a manner to ensure freedom from hazard.
- Avoid blocking the air intakes of the HoverTech Air Supply.



Avoid electric shock. Do not open air supply.

Note: Two versions of EMS Evacuation Hoverlack® Air Patient Lift



SEPARATE INFLATE/DEFLATE VALVES.

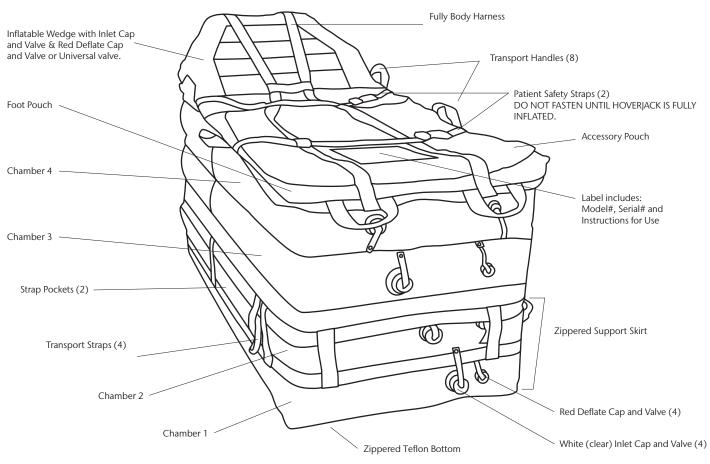
[Mfg before 6/2023]

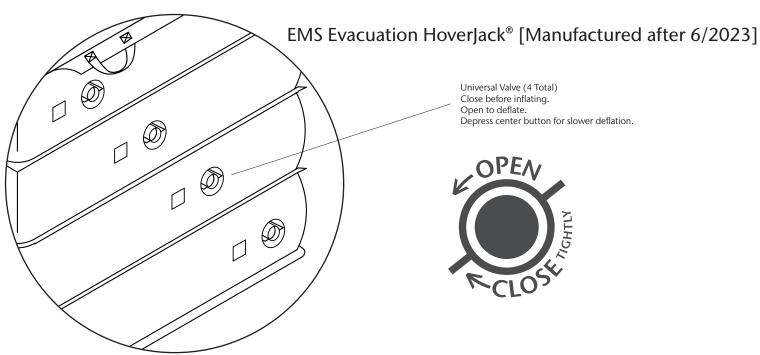


UNIVERSAL VALVES.
[Mfg after 6/2023]



Part Identification - EMS Evacuation HoverJack® [Manufactured before 6/2023]







EMS Evacuation Hoverlack® Instructions for Use as an Air Patient Lift

- 1. Place EMS Evacuation HoverJack® on the floor next to the patient, making sure the chamber with Valve #4 is on the top and the chamber with Valve #1 is against the floor.
- 2. Make certain that all four red-capped deflation valves are capped tightly or Universal Valves are closed BEFORE INFLATING.
- 3. Log roll patient onto the deflated EMS Evacuation HoverJack, and position patient with feet at the valve end where indicated. Alternatively, the patient can be placed on top of the EMS Evacuation HoverJack using the HoverMatt® Air Transfer System (see HoverMatt manual for instructions). If the HoverMatt is used, make certain that the HoverMatt and patient are properly centered on the EMS Evacuation Hoverlack. Always deflate the HoverMatt prior to inflating EMS Evacuation HoverJack.
- 4. Always use a minimum of two caregivers while using the EMS Evacuation HoverJack during inflation/transport.
- 5. One caregiver should remain on the side of the patient during inflation with the second caregiver managing inflation. If available, a third caregiver should attend to the other side of the patient.
- 6. Plug the HoverTech Air Supply power cord into an electrical outlet.
- 7. Caregiver at foot end turns on HoverTech Air Supply to initiate air flow.
- To begin inflation hold hose against valve #1 of EMS Evacuation HoverJack (valve closest to the floor).
- 9. When fully inflated, remove hose. Chamber remains inflated.
- 10. Secure patient safety straps.

- 11. Using the same process: move to valve #2, valve #3 and valve #4 in exact succession, or until the EMS Evacuation Hoverlack reaches desired height.
- 12. Turn off HoverTech Air Supply, and attach valve caps, if desired and applicable.
- 13. Move EMS Evacuation HoverJack to desired location.
- 14. If transferring from the EMS Evacuation HoverJack onto an adjacent surface, unbuckle patient safety straps.
- 15. If it is necessary to lower patient, slowly open the uppermost valve. When chamber is fully deflated, move in succession downwards to fully deflate. Depress center button for slower deflation.



CHAMBERS MUST BE FULLY INFLATED TO ENSURE STABILITY.



DO NOT RELEASE MULTIPLE CHAMBERS AT ONCE WHEN LOWERING A PATIENT.

EMS Evacuation Hoverlack® Instructions for Use as an Evacuation Device

- 1. To inflate, follow steps 1-11 of instructions for use as an air patient lift.
- Unzip foot pouch and position EMS Evacuation Hoverlack as close as possible to adjacent surface.
- 3. Transfer patient from bed or stretcher onto EMS Evacuation Hoverlack.
- 4. Ensure that the patient's body is centered, and zip foot pouch.
- 5. If necessary, inflate the head-end wedge.
- 6. Using the buckles, secure safety straps over the patient and adjust until taut.
- 7. Using foot end transport straps and handles located around the perimeter of the EMS Evacuation HoverJack, pull patient to the nearest stairwell, and position the EMS Evacuation Hoverlack so that the patient can be taken down the stairs feet first.
- 8. Before descending the stairs, chambers 3 and 4 must be fully deflated. To release air, slowly open valve #4. When chamber #4 is fully deflated, repeat the process for chamber #3. DO NOT RELEASE MULTIPLE CHAMBERS AT ONCE.
- 9. Retighten patient safety straps after deflating chamber #3 and #4 to ensure patient is secure.

- 10. Caregiver at head end will control evacuation using head end transport straps. Caregiver at head end should be physically stronger.
- 11. While 2 caregivers hold the headend transport straps and handles, the foot end caregiver will begin to pull the patient down the stairs with the foot end transport strap. The Teflon-infused bottom material will allow the EMS Evacuation HoverJack to slide down each flight. If necessary, the foot end caregiver can use their thigh to brace the EMS Evacuation HoverJack and slow the descent. Once at ground level, move the patient to safety.



WHEN LOWERING A PATIENT ENSURE THAT PATIENT REMAINS CENTERED ON THE EMS EVACUATION HOVERIACK. AND THAT THEIR HEAD DOES NOT SLUMP FORWARD AND RESTRICT BREATHING DURING EVACUATION.



A MINIMUM OF THREE CAREGIVERS (TWO AT HEAD END AND ONE AT FOOT END) SHOULD BE USED TO EVACUATE PATIENT DOWN STAIRS).



Product Specifications/Required Accessories

Material:	Topside Material: Nylon oxford / nylon Skirt: Cordura® fabric Underside Material: Teflon®	
Construction:	RF-Welded	
Width:	32" (81cm)	
Length:	72" (183 cm)	
Height:	30" (76 cm) Inflated [each chamber 7 1/2" (19 cm)]	

Model #: HJ32EV-2





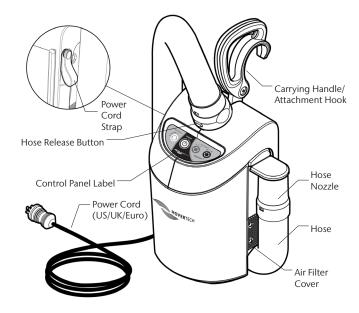
Weight limit: 1200 lbs (544 kg) for vertical lift. Weight limit: 700 lbs (318 kg) for stairwell evacuation.

REQUIRED ACCESSORY:

Model #: HTAIR1200 (North American Version) – 120V~, 60 Hz, 10A Model #: HTAIR2300 (European Version) – 230V~, 50 Hz, 6A Model #: HTAIR1000 (Japanese Version) – 100V~, 50/60 Hz, 12.5A Model #: HTAIR2356 (Korean Version) – 230V~, 50/60 Hz, 6A

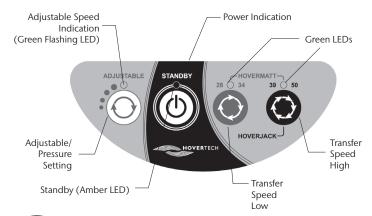
Model #: Air200G (800 W) (North American Version) – 120V~, 60 Hz, 10A Model #: Air400G (1100 W) (North American Version) – 120V~, 60 Hz, 10A

Part Identification – HT-Air® Air Supply



WARNING: The HT-Air is not compatible with DC power supplies. The HT-Air is not for use with the HoverJack Battery Cart.

HT-Air® Keypad Functions





ADJUSTABLE: For use with HoverTech air-assisted positioning devices. There are four different settings. Each press of the button increases the air pressure and rate of inflation. The Green Flashing LED will indicate the inflation speed by the number of flashes (i.e. two flashes equals the second inflation speed).

All of the settings in the ADJUSTABLE range are substantially lower than the HoverMatt and HoverJack settings. The ADJUSTABLE function is not to be used for transferring.

The ADJUSTABLE setting is a safety feature that can be used to ensure the patient is centered on HoverTech air-assisted devices and to gradually accustom a patient who is timid or in pain to both the sound and functionality of the inflated devices.



STANDBY: Used to stop inflation/air flow (Amber LED indicates STANDBY mode).



HOVERMATT 28/34: For use with 28" & 34" HoverMatts and HoverSlings.



HOVERMATT 39/50 & HOVERJACK: For use with 39" & 50" HoverMatts and HoverSlings and 32" & 39" HoverJacks.

Air200G/Air400G Air Supplies

If using HoverTech's Air200G or Air400G Air Supplies, press the grey button on the top of the canister to initiate air flow. Press the button again to stop air flow.



Cleaning & Maintenance



In between patient use, the EMS Evacuation Hoverlack should be wiped down with a cleaning solution used by your facility for medical equipment disinfection. A 10:1 bleach solution (10 parts water: one part bleach) or disinfectant wipes can also be used. NOTE: Cleaning with bleach solution may discolor fabric.

First remove any visible soil, then clean the area according to the cleaning product manufacturer's recommended dwell time and level of saturation. A gentle scrub brush can also be used on the affected area, if necessary, to help penetrate the EMS Evacuation Hoverlack material.



Do not launder the EMS Evacuation HoverJack.

PREVENTIVE MAINTENANCE

The EMS Evacuation HoverJack should be periodically inspected to ensure the following:

- All patient safety straps are attached.
- Transfer handles are all attached.
- All buckles and zippers are intact and operational.
- There are no punctures or tears.

EMS Evacuation Hoverlack with separate inflate/deflate valves:

- Inflate valves are all self-sealing with no evident leakage.
- All deflate valves are equipped with a red cap.
- The red caps are intact.

EMS Evacuation HoverJack with Universal Valves:

• Universal valves open fully and close tightly.

AIR SUPPLY CLEANING AND MAINTENANCE

See air supply manual for reference.

NOTE: CHECK YOUR LOCAL/STATE/FEDERAL/INTERNATIONAL **GUIDELINES BEFORE DISPOSAL.**

Transportation and Storage

This product does not require any special storage conditions



Returns and Repairs

All products being returned to HoverTech must have a Returned Goods Authorization (RGA) number issued by the company. Please call (800) 471-2776 and ask for a member of the RGA Team who will issue you an RGA number. Any product returned without an RGA number will cause a delay in the repair time.

Returned products should be sent to:

HoverTech Attn: RGA # 4482 Innovation Way Allentown, PA 18109

For European companies, send returned products to:



For product warranties, visit our website:

https://hovermatt.com/standard-product-warranty/



HoverTech

4482 Innovation Way Allentown, PA 18109

www.HoverMatt.com Info@HoverMatt.com

These products comply with the standards applicable for Class 1 products in the Medical Device Regulation (EU) 2017/745 on medical devices.



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In case an adverse event in relation to the device, incidents should be reported to our authorized representative. Our authorized representative will forward information to the manufacturer.



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