Hausted® by graham-field

ELECTRIC ALL-PURPOSE BARIATRIC STRETCHER 5B800ST

ELECTRIC ALL-PURPOSE BARIATRIC STRETCHER WITH NAVIGATOR SMART DRIVE SYSTEM 5B800ST-PD OPERATING MANUAL

SAVE THIS MANUAL FOR FUTURE USE.

HP_5B800ST-INS-LAB-RevC25

CONTENTS

INTRODUCTION — A WORD FROM GF HEALTH PRODUCTS, INC.

This manual contains important information on proper use and maintenance of the Hausted® 5B800 Series All-Purpose Bariatric Stretchers. All personnel involved in the use and maintenance of this equipment must carefully review and comply with the warnings, cautions and instructions contained in this manual. These instructions are important to protect the health and safety of personnel operating these models and should be retained in a conveniently accessible area for quick reference.

Complete instructions for uncrating and putting your new equipment in service, as well as equipment drawings, have been furnished. If missing, contact GF Health Products, Inc. ("GF") for replacement copies, giving the serial number and model number of the unit.

GF Health Products, Inc. carries a complete line of accessories for use with these stretchers; your representative will gladly review these with you.

Indications for Use

The Hausted 5B800 Series All-Purpose Bariatric Stretchers are intended for a healthcare professional to use in patient treatment, transport, or recovery. This product has an expected service life of five years.

Patient positioning is convenient and easy for both care provider and patient with the electric/ battery powered controls for adjustment of height, backrest, and knee-flex.

To ensure the basic safety of the patient, the stretcher is designed, tested, and evaluated to IEC Standard 60601-1 and in accordance with IEC 60601-2-46 wherein the essential performance in any single fault or combined fault condition is no unwanted movement of the stretcher when in use. See also page 7 – Electromagnetic Compatibility (EMC) information. The stretcher is tested and certified to IEC 60601-1-2 (Ed. 4) for EMC.

Service Information

A thorough preventive maintenance program is essential to safe and proper unit operation. This manual contains maintenance schedules and procedures which should be followed for satisfactory equipment performance.

We encourage you to contact GF Health Products, Inc. with maintenance concerns.

Advisory

A listing of the safety precautions to be observed when operating and servicing this equipment can be found in Section 1 of this manual. Do not operate or service the equipment until you have become familiar with this information. Any alteration of this equipment not authorized or performed by GF Health Products, Inc., could affect its operation, will void the warranty, could violate national, state, and local regulations, and could jeopardize your insurance coverage.

Info: *Column 1 below applies only if product was purchased outside the U.S.*

CE EU REP

EU Authorized Representative:

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1.770.368.4700 Main
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www.Hausted.com

Class 1 Equipment

Type B Equipment

Equipment not suitable for use in the presence of flammable anesthetic mixture with air or oxygen or nitrous oxide.

IPX4 Not suitable for continuous operation (Duty Cycle: 10% 2 Min. in 18 Min.)

Info: *The base language of this document is ENGLISH. Any translations must be made from the base language document.*

1 LIST OF WARNINGS AND CAUTIONS

⚠ **IMPORTANT: Before using the All-Purpose Bariatric Stretcher, please read and adhere to the following safety precautions and warnings. Failure to do so could result in serious personal injury or damage to the Stretcher.**

Always consult your healthcare professional to determine safe methods most suitable for your individual abilities. Protect yourself, your attendant, and the All-Purpose Bariatric Stretcher by having it serviced regularly. If you experience any malfunction, contact your Graham-Field authorized distributor immediately, as a hazardous condition could result, causing personal injury or damage to the Stretcher.

Periodic inspection, adjustment and replacement of worn parts are necessary to provide years of excellent service. Maintenance MUST be performed by qualified personnel ONLY.

SAVE THESE INSTRUCTIONS.

SIGNIFICANCE OF SAFETY STATEMENTS

Please note the following special statements, used throughout this manual, and their significance:

⚠ **DANGER: Indicates a potential hazard situation or unsafe practice that, if not avoided, will result in death or serious personal injury.**

⚠ **WARNING: Indicates a potential hazard situation or unsafe practice that, if not avoided, could result in death or serious personal injury.**

⚠ **CAUTION: Indicates a potential hazard situation or unsafe practice that, if not avoided, could result in minor or moderate personal injury.**

▲ **NOTICE: Indicates a potential hazard situation or unsafe practice that, if not avoided, could result in product or property damage.**

Info: *Provides application recommendations or other useful information to ensure that you get the most from your product.*

DANGER / WARNING / CAUTION / NOTICE Summary

The following is a listing of the safety precautions which must be observed when operating and servicing this equipment. These precautions are repeated (in whole or in part), where applicable, throughout the manual.

DANGER: To Reduce the Risk of Burns, Fire, or Electric Shock

⚠ **DANGER: SHOCK HAZARD — To reduce the risk of electric shock, unit is to be serviced by qualified personnel only.**

⚠ **DANGER: SHOCK HAZARD — To avoid the risk of electric shock, this equipment must only be connected to supply mains with protective earth.**

WARNING: To Reduce the Risk of Personal Injury

⚠ **WARNING: LACERATION HAZARD — When cutting bands ALWAYS use a tool specifically designed for that purpose. This will help to avoid personal injuries frequently incurred when bands are cut and tension released.**

⚠ **WARNING: DO NOT sit on end — tipping may occur.**

⚠ **WARNING: Ensure IV rod is inserted completely into socket up to the arrow BEFORE applying load.**

WARNING — CAUTIONS AND PROPER OPERATION

⚠ **WARNING: The Series 5B800 stretchers have a maximum patient weight capacity of 800 lb (363 kg), EVENLY DISTRIBUTED.**

⚠ **WARNING: The Series 5B800 stretchers have a maximum weight, including equipment weight, and patient weight of 1200 lb (544 kg), EVENLY DISTRIBUTED.**

⚠ **WARNING: Patient entry, egress and transfer should ALWAYS be performed with the brakes locked.**

⚠ **WARNING: The patient transport position is pushing from the patient HEAD END in the supine position.**

⚠ **WARNING: The brakes should ALWAYS be locked and patient side rails up when patient is not in transport.**

⚠ **WARNING: Clip patient pendant to rail when not in use – keep cord clear of moving parts.**

⚠ **WARNING: This electric powered stretcher is equipped with a built-in battery backup system, but the unit should remain plugged into wall receptacle during normal use. The battery backup is intended for transport and EMERGENCY ONLY.**

⚠ **WARNING: The stretcher has a warning label on both the head and foot end stating: "Do not sit on end - as tipping may occur."**

⚠ **WARNING: Patient entry, egress, and transfer from the stretcher should always be from the center side rail location with the side rail in the down position and brakes locked.**

⚠ **WARNING: The back quick drop handles are intended to be used during emergency situations only.**

⚠ **WARNING: To turn on electric controls, plug into wall receptacle. To turn off, remove plug from wall receptacle. Electric powered stretchers do not have a separate on / off switch.**

⚠ **WARNING: ALWAYS disconnect the power source whenever troubleshooting or servicing any electric powered stretcher.**

⚠ **WARNING: Cables can become pinched. Keep cables away from column.**

⚠ **WARNING: Steam cleaning and pressure washing of stretcher is not recommended and can void warranty.**

⚠ **WARNING: DO NOT modify the equipment without the authorization of the manufacturer.**

⚠ **WARNING: When lowering the rails, ensure patient and caregiver body and extremities are clear of pinch points before operating the rail.**

⚠ **WARNING: Keep hands clear of pinch points.**

⚠ **WARNING: Stow away power cord when not in use to prevent injury or damage.**

ELECTROMAGNETIC COMPATIBILITY (EMC) INFORMATION

⚠ **WARNING: Medical Electrical Equipment needs special precautions regarding EMC and needs to be installed and put into service according to the EMC information provided in this manual.**

⚠ **WARNING: Electronic equipment may be influenced by Radio Frequency (RFI). Caution should be exercised with regard to the use of portable communications in the area around such equipment. Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 12 inches (30 cm) to any part of the Hausted equipment including specified Hausted equipment cables. Degradation of the performance of the Hausted equipment could result.**

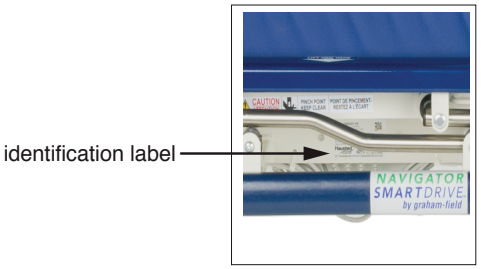
⚠ **WARNING: If RFI causes erratic behavior, unplug the electric Hausted equipment immediately. Leave unplugged while transmission is in progress.**

⚠ **WARNING: The use of accessories, transducers, and cables other than those specified by the manufacturer may result in increased emissions or decreased immunity of the Hausted equipment. GF cables and accessories include motor cables, mains cable, pendant cables, and back up battery and cable.**

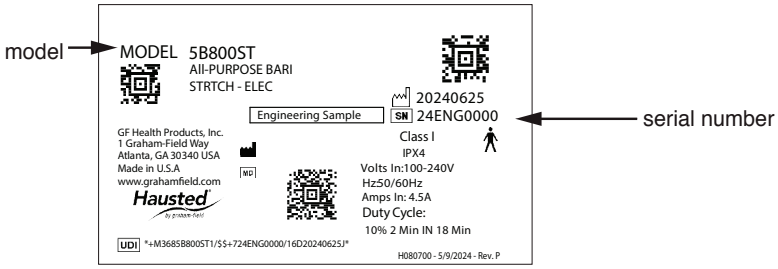
⚠ **WARNING: This equipment should NOT BE used adjacent to or stacked with other equipment. If adjacent or stacked use with other equipment is necessary, this Hausted equipment and the other equipment should be observed to verify that they are operating normally.**

⚠ **WARNING: The EMISSIONS characteristics of this equipment make it suitable for use in industrial areas and hospitals (CISPR 11 class A). If it is used in a residential environment (for which CISPR 11 class B is usually required) this equipment might not offer adequate protection to radio frequency communication services. The user might need to take mitigation measures, such as relocating or reorienting the equipment.**

IDENTIFICATION LABEL



Identification label location



Identification label example

The stretcher identification label identifies the stretcher serial number and model, essential information when ordering replacement parts or claiming parts under warranty. The identification label, shown above, is beneath the patient left surface at midpoint of frame. Have this information ready when calling our Customer Service or Technical Support staff at 1.770.368.4700; it will allow us to better assist you and quickly answer your questions and concerns.

2 UNCRATING INSTRUCTIONS

IMPORTANT — REPORT ANY SHIPPING DAMAGE IMMEDIATELY:

⚠ **WARNING: Inform shipper of any damage — leave carton intact. Leave equipment in the receiving area until inspection is complete.**

NOTICE — POSSIBLE EQUIPMENT DAMAGE:

▲ **NOTICE: The crate contains fragile, expensive medical equipment. Uncrate and handle carefully. If after uncrating the equipment you find any damage (no matter how slight), report the damage to GF Health Products, Inc.**

WARNING — PERSONAL INJURY HAZARD:

⚠ **WARNING: When cutting bands, always use tool specifically designed for that purpose. This will help avoid personal injuries possibly incurred when bands are cut and tension is released.**

ENVIRONMENTAL CONDITIONS

Operating	
Temperature	5°C to 40°C
Relative Humidity	20% to 90% @ 86°F (30°C)
Atmospheric Pressure	700 to 1060 hPa

Storage and Transport	
Temperature	-10°C to 50°C
Relative Humidity	20% to 90% @ 86°F (30°C)
Atmospheric Pressure	700 to 1060 hPa

UNPACKING INSTRUCTIONS

IMPORTANT: Follow each step in the order shown in these instructions.

Your Hausted equipment has been carefully packed at our manufacturing plant to ensure safe shipment to your medical facility. There are several procedures you must follow to put your new equipment in service. These procedures only take a few minutes to complete and all are required to ensure proper operation of the equipment.

1. Cut the two bands around the shipping carton.
2. Remove the top half of the carton and cut one side of the bottom half.
3. Ensure the fifth wheel / power drive wheel clears the edge of the skid when removing product (where applicable), then remove the equipment from the carton.
4. Check to see if all features of the equipment work properly. If all the features work, advance to step 5. If any features do not work properly, call GF Health Products, Inc. at 1.770.368.4700.

Info: *Plug the unit into a wall socket prior to checking any electric features. The battery will reach full charge after approximately 10 hours.*

5. Clean the equipment using mild detergent to remove any dirt accumulated during shipment, and place the equipment into service.

3 OPERATING INSTRUCTIONS

3.1 5B800 STRETCHER SPECIFICATIONS

Info: *All dimensions, unless otherwise specified, are in inches and ±.375 (1 cm).*
GF Health Products, Inc. reserves the right to change specifications without notice.

Model	Bariatric 5B800ST & 5B800ST-PD
Height Range: High	40 in. ±1 in. [101.6 cm ±2.5 cm]
Height Range: Low	22 in. ±1 in. [55.9 cm] ±2.5 cm
Overall Width	37 in. [94 cm]
Overall Length	88.4 in. [224.5 cm]
Patient Surface Width	30 in. [76.2 cm]
Patient Surface Length	76.4 in. [194.1 cm]
Backrest Degree of Movement	0° - 85° ±2°
Knee Flex Degree of Movement	0° - 60° ±2°
Trendelenburg / Reverse Trendelenburg	15° ±2°
Retracto Rail Height	14 in. [35.6 cm]
Retracto Rail Length	63 in. [160 cm]
Casters	5.9 in. [15 cm] Electric Lock / Steer
Maximum Weight Capacity (EVENLY DISTRIBUTED)	800 lb [363 kg]
Applied Parts:	Pads, Linens, Side Rails, Some Accessories
Mattress Pad Thickness	5 in. [13 cm]

Electrical Specifications	
Product Classification:	1
Input Voltage:	100V - 240V ~ 50/60 Hz
Amperage:	Max. 4.0A
Duty Cycle:	10% (2 Min. in 18 Min.)
IP Rating:	IPX4 All Models
Grounding Protection:	Type B

3.2 FEATURES, WARNINGS AND PROPER OPERATION OPERATING INSTRUCTIONS

WARNINGS — CAUTIONS AND PROPER OPERATION (See Diagram Below)

A. ⚠ WARNING: DO NOT sit on end — tipping may occur.

B. ⚠ WARNING: The stretchers have a warning label located at the Head-End (Back Side of Back Section) stating: Maximum patient weight 800 LB (363 kg) for 5B800 models.

C. ⚠ WARNING: Patient entry, egress and transfer should always be done with the brakes locked.

D. ⚠ WARNING: The brakes should always be locked and patient side rails up when patient is not in transport.

E. ⚠ WARNING: The patient pendant has a warning label stating: Clip pendant to rail when not in use — keep cord clear of moving parts. As is stated on the cable adjacent to the pendant.

F. ⚠ WARNING: All electric stretchers are equipped with a built-in battery backup system, but it is recommended that the unit remain plugged into wall receptacle during normal use. The battery backup is recommended for transport and EMERGENCY only.

G. ⚠ WARNING: The back quick drop handles are intended to be used to lower a patient during EMERGENCY situations only.

H. ⚠ WARNING: Ensure rail is locked before leaving patient.

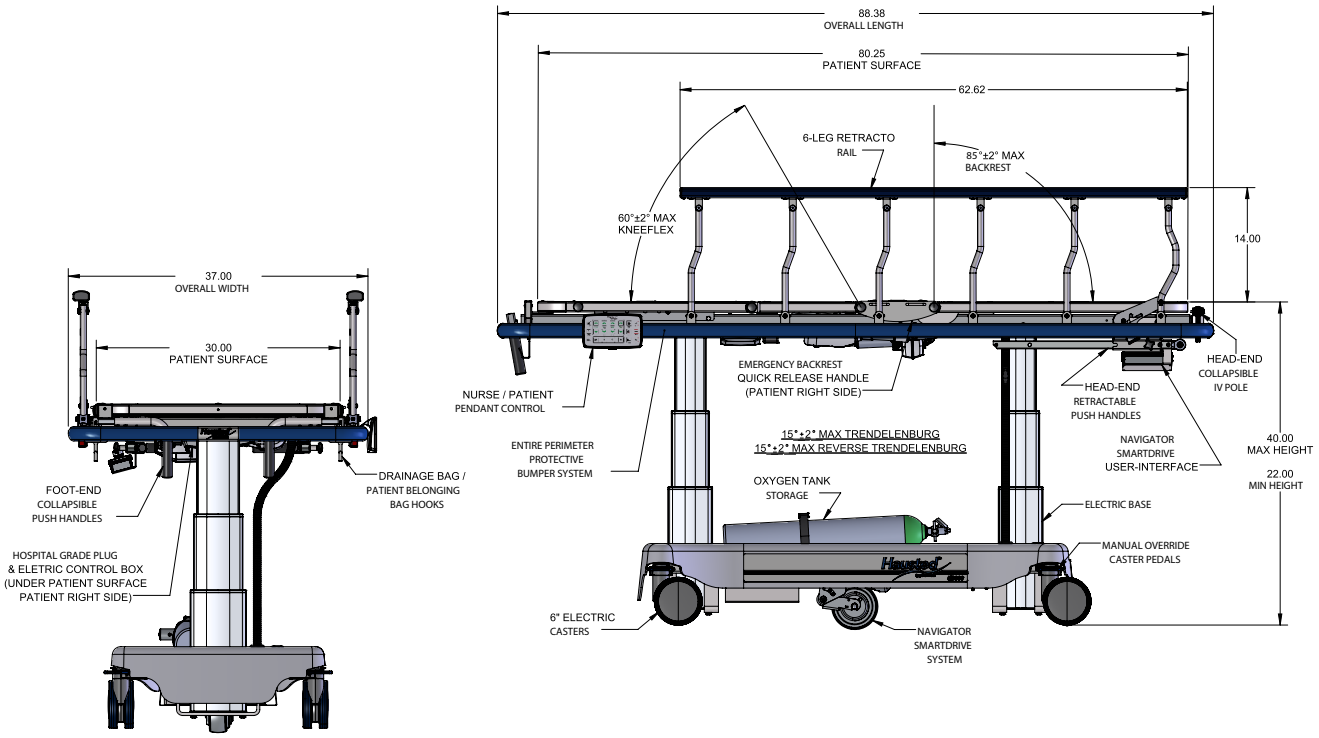
I. ⚠ WARNING: When lowering the rails, ensure patient and caregiver (or attendant) body and extremities are clear of pinch points before operating the rail.

J. ⚠ WARNING: To turn on electric controls, plug into wall receptacle; to turn off, remove plug from wall receptacle. The electric powered stretchers do not have a separate on / off switch.

K. ⚠ WARNING: Always disconnect the power source whenever servicing any electric powered stretcher.

L. ⚠ WARNING: Keep hands clear of pinch points.

M. ⚠ WARNING: Stow away power cord when not in use to prevent injury or damage.



WARNINGS — CAUTIONS AND PROPER OPERATION (See List on Previous Page)

3.3 BRAKING AND STEERING OPERATION WITH SMART CASTER TECHNOLOGY

3.3.1 Applying the Brakes

To apply the four-wheel braking system with the pendant, press the pendant Brake On / Off button; the LED above the button then illuminates green (Figure 3.3-1), an audible beep will be heard, and the blue caster pedals on all four corners of the stretcher automatically lower to brake position (Pedal Down) (Figure 3.3-2), and all four caster wheels will then be locked from swiveling and rotating.

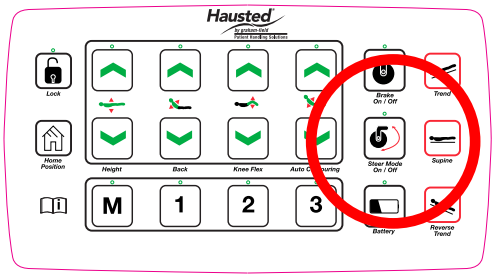


Figure 3.3-1

Info: *To prevent unintended movement, the stretcher is equipped with an Automatic Braking Feature and a Brake Alarm System. The brakes will engage automatically after the unit has been stationary for 3 consecutive minutes. The Brake Alarm will sound (5 quick beeps) at 5 seconds, before the brakes engage to provide warning that the casters have locked.*

Info: *An audible beep will be heard with each change of caster position (Brake / Neutral / Steer-Lock).*

To apply the four-wheel braking system manually, gently depress the blue caster pedal at any of the four corners of the stretcher until the pedal stops (Figure 3.3-2); the LED above the Brake On / Off button then illuminates green (Figure 3.3-1), an audible beep will be heard, and the remaining three pedals will then automatically lower to brake position (Pedal Down) (Figure 3.3-2), and all four caster wheels will then be locked from swiveling and rotating.

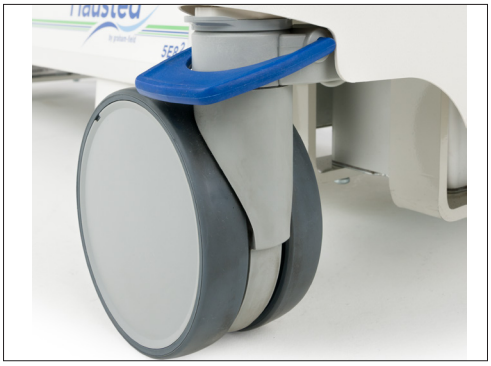


Figure 3.3-2

▲ NOTICE: DO NOT apply excessive force to pedal when manually applying brakes.

3.3.2 Unlocking the Brakes

To unlock the brakes with the pendant, press the pendant ***Brake On / Off*** button; the LED above the button then goes out (Figure 3.3-3), an audible beep will be heard, and the blue caster pedals on all four corners of the stretcher will then automatically rise to neutral position (Pedal Horizontal) (Figure 3.3-4), and all four caster wheels will then rotate and swivel freely.

To unlock the brakes manually, gently lift the blue caster pedal at any of the four corners of the stretcher until the pedal stops (Figure 3.3-6); the LED above the Brake On / Off button then goes out (Figure 3.3-3), an audible beep will be heard, and the remaining three pedals will then automatically rise to neutral, and all four caster wheels will then rotate and swivel freely.

⚠ **WARNING: Ensure the brakes are locked when the patient is not being transported.**

▲ **NOTICE: DO NOT apply excessive force to pedal when manually unlocking brakes.**

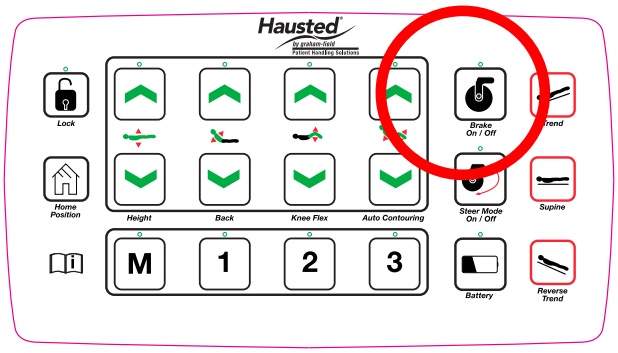


Figure 3.3-3

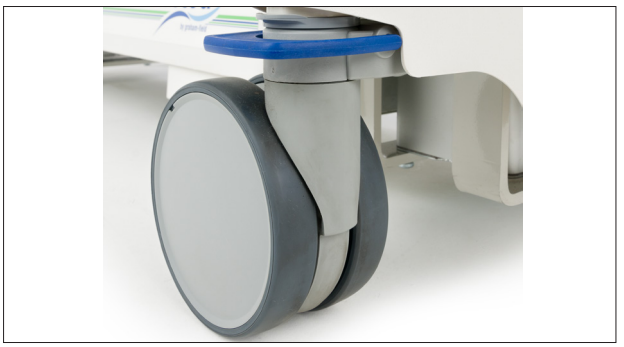


Figure 3.3-4

3.3.3 Activating Steer Mode — Pendant

To activate Steer Mode with the pendant, press the pendant ***Steer Mode On / Off*** button. The LED above the button then illuminates green (Figure 3.3-5), an audible beep will be heard, and the blue caster pedals at the patient foot-end of the stretcher will automatically rise to Steer-Lock position (Pedal Up) (Figure 3.3-6).

The patient head-end caster pedals will automatically rise to neutral position (Pedal Horizontal) (Figure 3.3-4), and the stretcher will be ready for transport.
From the patient head-end, push the stretcher forward – both front casters will lock into Steer-Lock position.

The stretcher will steer along a straight path, maneuver corners, and change direction with minimal effort.

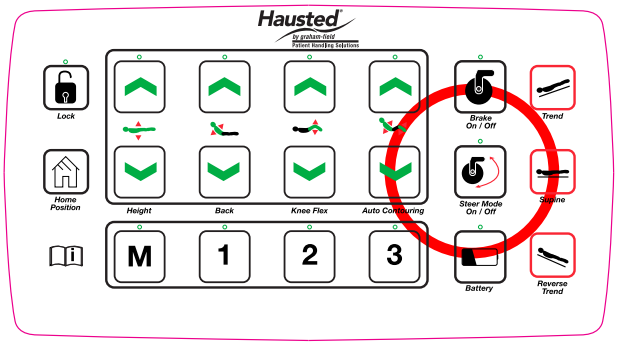


Figure 3.3-5

NOTE: ***The steer mode function (both by pendant or manually) is disabled for stretchers with the Navigator Smart Drive System. Activation with the pendant results in an audible beep. Manual activation will immediately return to the neutral position.***

Info: *The casters will lock into Steer-Lock position when turned to 6 o'clock or 12 o'clock, with the stretchers' patient foot-end being 12 o'clock.*

Info: *Depending upon how the casters are oriented when they lock, they can lock into Steer-Lock position while trailing (6 o'clock) or Leading (12 o'clock). If the casters lock in the opposite orientations, steering may be more difficult.*

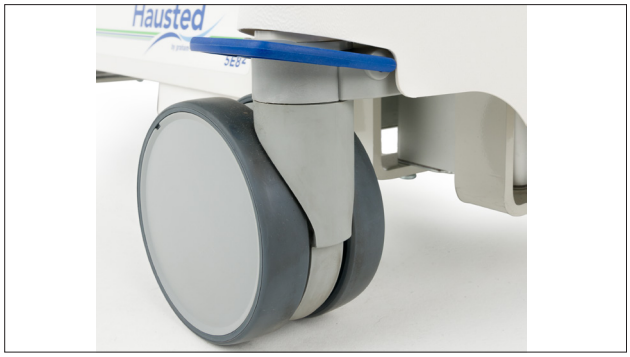


Figure 3.3-6

Info: *The pendant will only lock the patient foot-end casters into Steer-Lock Position, which is ideal for pushing the stretcher from the patient head-end.*

3.3.4 Activating Steer Mode – Manually

NOTE: ***All four casters must be in neutral position (Pedal horizontal) (see Figure 3.3-4 on page 12) before manually activating Steer Mode.***

When pushing from the head end: Make sure all four casters are in neutral position (Pedal Horizontal) (see Figure 3.3-4 on page 12); activate foot-end Steer Mode by lifting the blue pedal upward on either head-end caster until the pedal stops (see Figure 3.3-6 on page 12). Both foot-end pedals will rise into Steer-Lock position (Pedal Up) (see Figure 3.3-6 on page 12), the head-end pedals will remain in neutral position (Pedal Horizontal) (see Figure 3.3-4 on page 12), the LED above the pendant ***Steer Mode On / Off*** button will illuminate green (see Figure 3.3-5 on page 12), and an audible beep will be heard. From the patient head-end, push the stretcher forward – both front casters will lock into Steer-Lock position and the stretcher will be ready for transport.

When pushing from the foot end: Make sure all four casters are in neutral position (Pedal Horizontal) (see Figure 3.3-4 on page 12); activate head-end Steer Mode by lifting the blue pedal upward on either foot-end caster until the pedal stops (see Figure 3.3-6 on page 12). Both head-end pedals will rise into Steer-Lock position (Pedal Up) (see Figure 3.3-6 on page 12), the foot-end pedals will remain in neutral (Pedal Horizontal) (see Figure 3.3-4 on page 12), the LED above the pendant ***Steer Mode On / Off*** button will illuminate green (see Figure 3.3-5 on page 12), and an audible beep will be heard. From the patient foot-end, push the stretcher forward – both front casters will lock into Steer-Lock position and the stretcher will be ready for transport. Info: It is not possible to lock the head end casters into Steer-Lock position with the pendant.

This can only be done with the foot end manual activation.

Info: *It is not possible to lock the head-end casters into Steer-Lock position with the pendant. This can only be done with the foot-end manual activation.*

3.3.5 Deactivating Steer Mode — Pendant

To deactivate Steer Mode with the pendant, press the pendant ***Steer Mode On / Off*** button. The LED above the button will go out (Figure 3.3-7), and an audible beep will be heard, and the blue casters pedals in Steer-Lock position will automatically lower to neutral (Pedal Horizontal) (see Figure 3.3-4 on page 12). All four casters will now rotate and swivel freely.

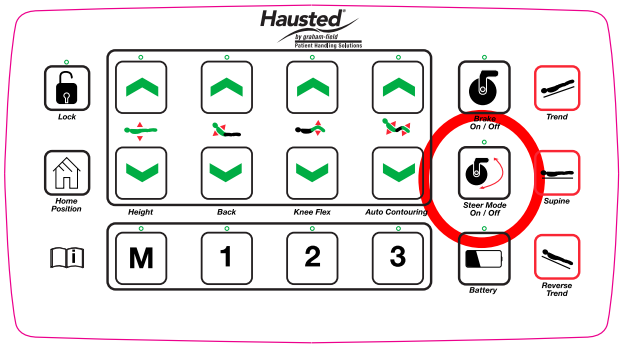


Figure 3.3-7

Info: *All four casters must be parallel to each other in the 6 o'clock or 12 o'clock position to be able to properly go into unlock / neutral position. Failure to place casters in this orientation may cause the Steer Mode On / Off button to not deactivate Steer Mode.*

3.3.6 Deactivating Steer Mode – Manually

Depress the blue caster pedal down to the neutral position (Pedal Horizontal) (see Figure 3.3-4 on page 12) on any caster locked in Steer-Lock position. The LED above the Steer Mode On / Off button will go out (see Figure 3.3-7 on page 13), an audible beep will be heard, and the other caster in Steer-Lock position will automatically lower to neutral position (Pedal Horizontal) (see Figure 3.3-4 on page 12). All four casters will now rotate and swivel freely. Depressing the blue pedal down past neutral until it stops will apply all four casters brakes (see Figure 3.3-2 on page 11), locking all four casters into brake position (Pedal Down).

Info: *After thirty seconds of no movement, Steer Mode automatically deactivates and all four caster wheels lock into brake position.*

⚠ **WARNING: To prevent unintended movement, activate or deactivate Steer Mode ONLY while the stretcher is stopped.**

Caster Pedal Positions

The table below shows the blue caster pedal in all three positions.

CASTER PEDAL POSITIONS		
	Pedal Up	Steer-Lock Position
	Pedal Horizontal	Neutral Position (Swivel)
	Pedal Down	Brake Position (Locked)

3.4 ELECTRIC CONTROL LOCATIONS

3.4.1 Pendant Control Storage Location

The pendant is located on the bumper rail on either side of the stretcher (Figure 3.4-1).

Figure 3.4-1

▲ NOTICE: Place pendant on bumper rail when not in use. Keep cord clear of moving parts.

3.4.2 Plug Location

This stretcher is equipped with a battery backup for transport but the unit should be plugged into a wall receptacle when not in transport. The plug is located on the patient right side of the stretcher (Figure 3.4-2). Do not position the unit so that it is difficult to disconnect the plug.

Figure 3.4-2

⚠ WARNING: The All-Purpose Bariatric Stretcher is equipped with a built-in battery backup system: nevertheless, the unit should remain plugged into wall receptacle during normal use. The battery backup is recommended for transport and emergency only.

3.4.3 Low Battery Alarm

This stretcher is equipped with an audible and visual low battery alarm. When the system requires charging, a continuous beep will sound during motor operation, the pendant LED above the ***BATTERY*** button will illuminate green, and the control box LED will illuminate amber.

3.4.4 Foot Control

Large red circle indicates optional foot control storage location (Figure 3.4-3). Small red circle indicates foot control plug-in location (Figure 3.4-3), which can also be used as an alternate pendant plug-in location.

Figure 3.4-3

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

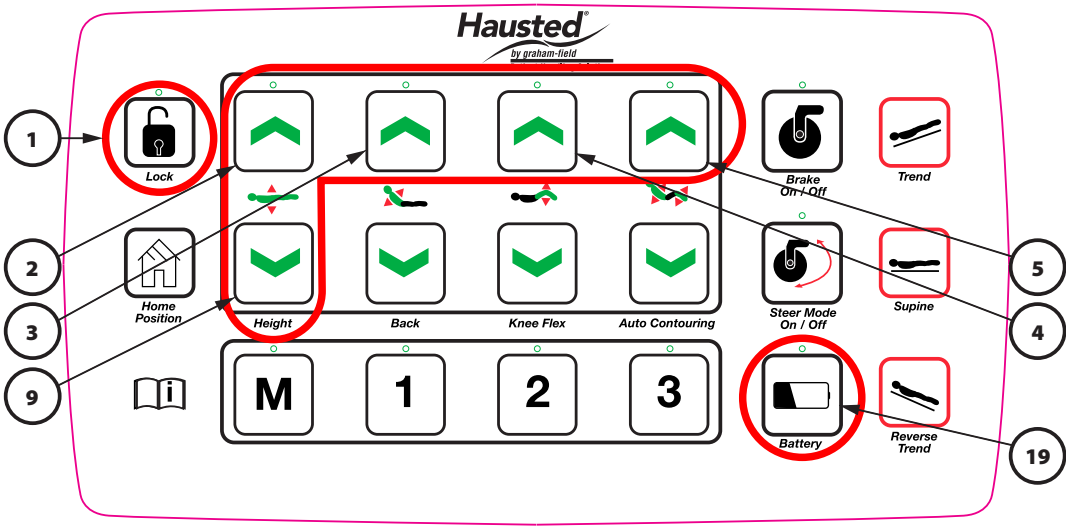
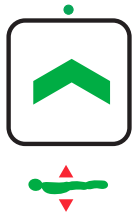


Figure 3.5-1

3.5.1 LOCK / UNLOCK (button 1)

1	 Lock  Battery	LOCK	Press and hold LOCK button (1) for three seconds to lock all functions. After five LED flashes, all four UP LED's (2-5) illuminate steady green, indicating they are now locked; an audible signal also indicates when locked and beeps up to three times until button is released.
		UNLOCK	Press and hold LOCK and BATTERY buttons (1 and 19) at the same time to unlock all functions. All UP-LED's will flash with three audible beeps indicating unlocked. All UP-LED's will continue to flash until BOTH buttons are released.
		UNLOCK INDIVIDUAL FUNCTION (PATIENT MODE)	Press and hold LOCK button (1) and press each UP button (2-5) to unlock each function individually. As each button is released, its LED will go out, indicating that function is unlocked.

3.5.2 HEIGHT (HI / LO) (buttons 2 and 9)

2		HEIGHT UP	Press and hold HEIGHT button (2) until desired height is achieved. LED illuminates steady green while pressed, goes out when released.
9	Height	HEIGHT DOWN	Press and hold HEIGHT DOWN button (9) until desired height is achieved. LED illuminates steady green while pressed, goes out when released.

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

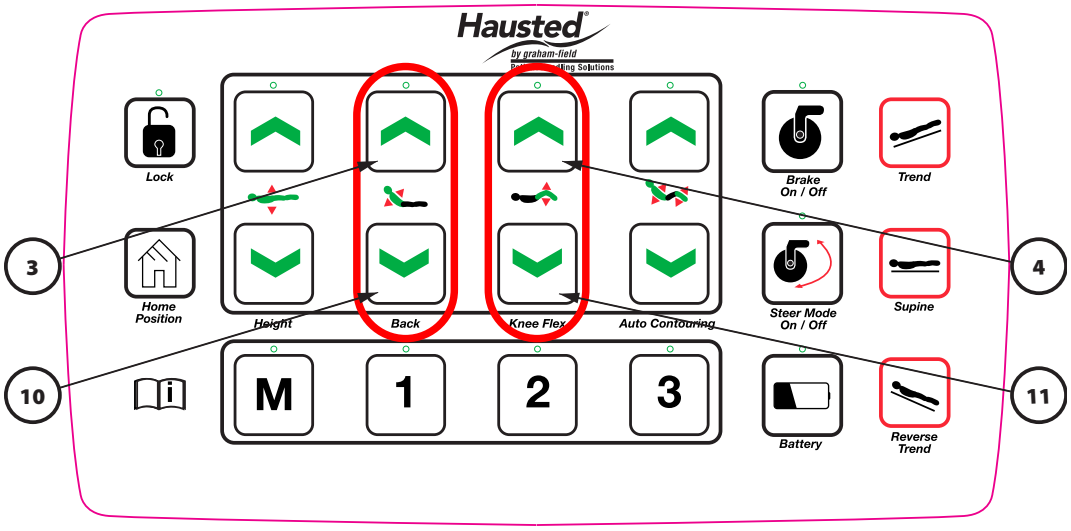
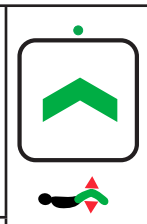


Figure 3.5-2

3.5.3 BACK SECTION UP / DOWN (buttons 3 and 10)

3		BACK UP	Press and hold BACK UP button (3) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.
10		BACK DOWN	Press and hold BACK DOWN button (10) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.

3.5.4 KNEE FLEX UP / DOWN (buttons 4 and 11)

4		KNEE FLEX UP	Press and hold KNEE FLEX UP button (4) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.
11		KNEE FLEX DOWN	Press and hold KNEE FLEX DOWN button (11) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

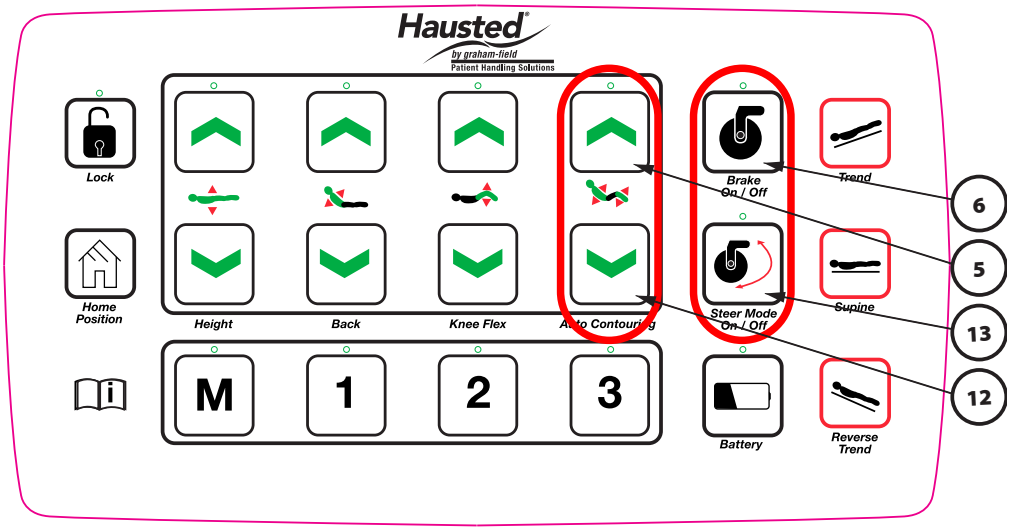


Figure 3.5-3

3.5.5 AUTO CONTOURING UP / DOWN (buttons 5 and 12)

5		AUTO CONTOURING UP	Press and hold AUTO CONTOURING UP button (5) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.
12		AUTO CONTOURING DOWN	Press and hold AUTO CONTOURING DOWN button (12) until desired position is achieved. LED illuminates steady green while pressed, goes out when released.

3.5.6 BRAKE and STEER MODE Operation (buttons 6 and 13)

6		BRAKE	Press BRAKE button (6) to toggle on / off. When brakes are locked (on), LED illuminates steady green; when brakes are unlocked (off), LED goes out. To prevent unintended movement, brakes lock automatically after stretcher is stationary for 3 consecutive minutes. Before brakes engage, a Brake Alarm will sound (5 quick beeps) 5 seconds before.
13		STEER MODE	Press STEER MODE button (13) to toggle on / off. When steer mode is activated, LED illuminates steady green; when steer mode is off, LED goes out. *Note: Steer mode is not operable when brake is activated.* *Note: Steer mode is disabled on stretchers equipped with the Navigator Smart Drive System.* *Note: Steer Mode button only locks the patient foot-end casters into Steer-Lock Position.*

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

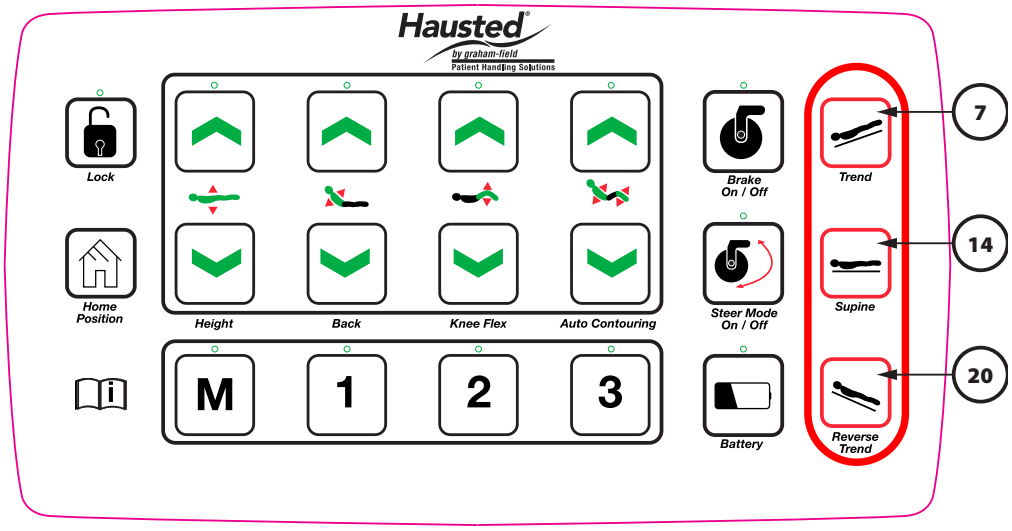


Figure 3.5-4A

3.5.7 TRENDELENBURG, SUPINE and REVERSE TRENDELENBURG Positions (buttons 7, 14 and 20)

7	 Trend	TRENDELENBURG	Press and hold TRENDELENBURG button (7) until desired position is achieved. The stretcher automatically lowers the back and knee-flex sections, tilts backward, and adjusts height. No LED.
14	 Supine	SUPINE	Press and hold SUPINE button (14) until desired position is achieved. The stretcher automatically levels the back and knee-flex sections and adjusts height. No LED.
20	 Reverse Trend	REVERSE TRENDELENBURG	Press and hold REVERSE TRENDELENBURG button (20) until desired position is achieved. The stretcher automatically lowers the back and knee-flex sections, tilts forward, and adjusts height. No LED.

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

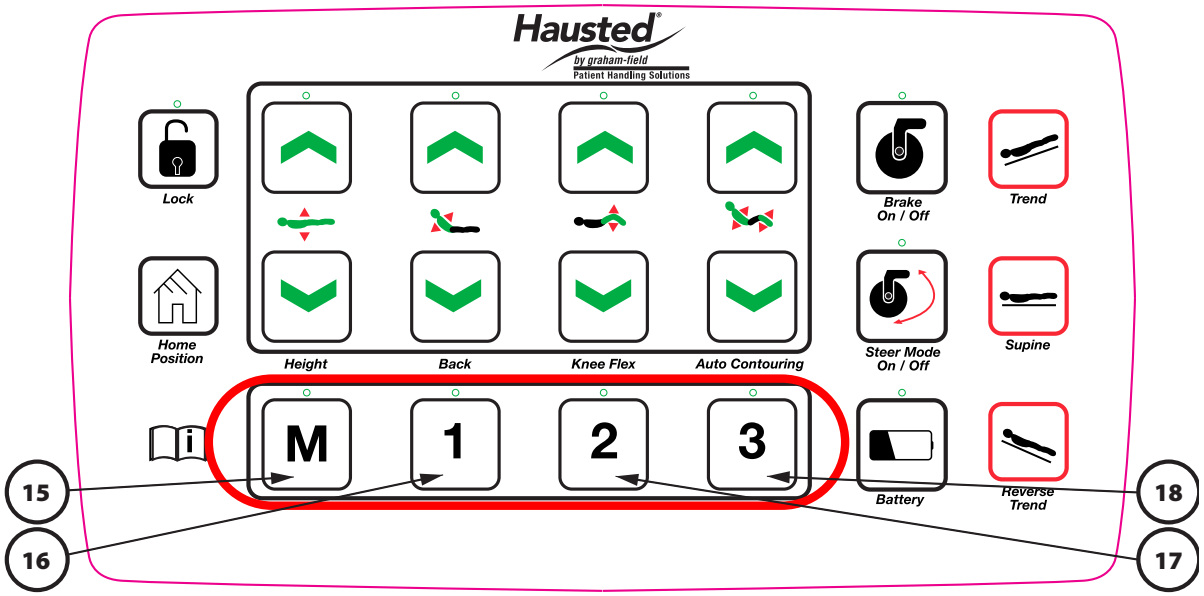
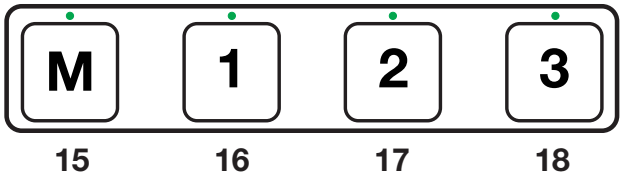


Figure 3.5-4B

3.5.8 Memory and Preset Functions (Buttons 15-18)



15	MEMORY	Position stretcher to desired position. Enter MEMORY MODE by pressing and holding MEMORY button (15) for three seconds until LED flashes. Once MEMORY LED flashes, simultaneously press and hold MEMORY button (15) and desired PRESET button (16, 17, or 18) until beeping stops (MEMORY LED will stop flashing and go out and PRESET LED will illuminate). Once MEMORY button and PRESET button are released, the position saves, LEDs go out, and MEMORY MODE exits. (After entering MEMORY MODE, PRESET buttons that illuminate are already programmed, but can be overwritten; PRESET buttons that don't illuminate are not yet programmed.)
16	PRESET 1	Press and hold PRESET 1 button until saved pre-programmed position is achieved. LED illuminates steady green while pressed, goes out when released.
17	PRESET 2	Press and hold PRESET 2 button until saved pre-programmed position is achieved. LED illuminates steady green while pressed, goes out when released.
18	PRESET 3	Press and hold PRESET 3 button until saved pre-programmed position is achieved. LED illuminates steady green while pressed, goes out when released.

3.5 HEIGHT AND PATIENT SURFACE ADJUSTMENT

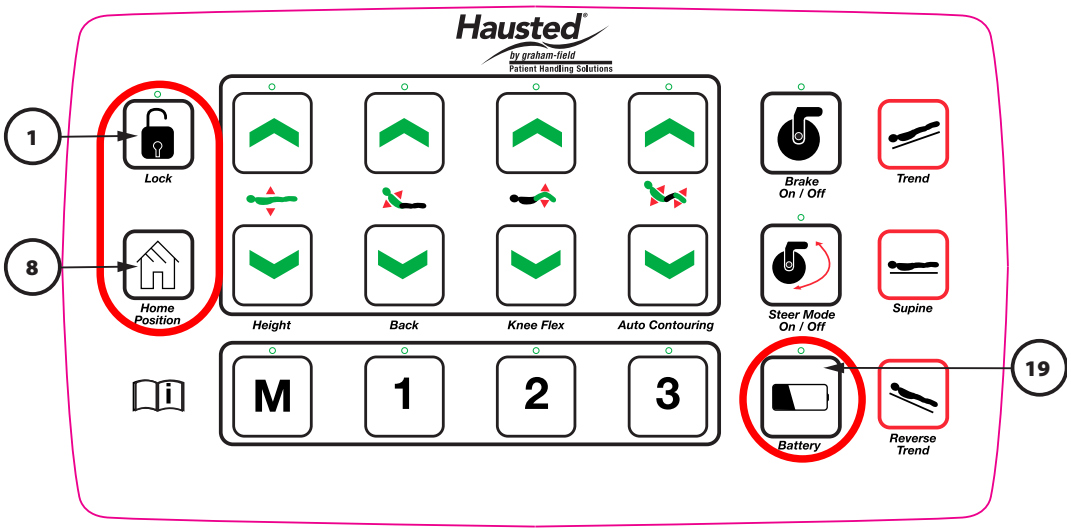


Figure 3.5-5

3.5.9 BATTERY (button 19)

19		BATTERY	LED illuminates steady green when battery discharges to 20% capacity or less. Stretcher will emit audible beeping sound when using the unit. Press and hold LOCK and BATTERY buttons (1 and 19) at the same time to unlock all functions. All UP-LED's will flash with three audible beeps indicating unlocked. All UP-LED's will continue to flash until BOTH buttons are released.

3.5.10 HOME (button 8)

8		HOME	Press and hold HOME button (8). The stretcher automatically lowers the back, and knee-flex sections and adjusts to lowest height. No LED.

3.5.11 Emergency Drop Back

This stretcher is equipped with a manual override function for the back section. This option should only be used in an emergency situation.

To activate the emergency drop back, support the back section, and pull outward on either red activating handles located on the patient right and left side under the seat to the rear (Figure 3.5-6).

Figure 3.5-6

Info: *When activating the emergency drop back, depending on the back angle, you may need to push down on the back in order to initiate the movement.*

Info: *When activating the emergency drop back, depending on how far the red activating handle is pulled outward, it is possible to control the descend speed of the back section.*

3.6 PUSH HANDLES

3.6.1 Operating the Head-End & Foot-End Push Handles

Info: *Push Handle operation is the same for both the Head-End & Foot-end of the stretcher.*

Raising the Push Handles: To use the Push Handle, hinge the push handle up/outward (Figure 3.6-2) unit it stops. Slide the push handle down into the handle bracket until it stops. Repeat process for handle on the other side (Figure 3.6-3).

Lowering the Push Handles: To stow away Push Handle, lift push handle up and out of bracket, then rotate (Figure 3.6-2) the push handle down/inward (Figure 3.6-1). Repeat process for other side.

⚠ **WARNING: To prevent unintended interference, ensure push handles are stowed away (as shown in Figure 3.6-1) when not in use.**

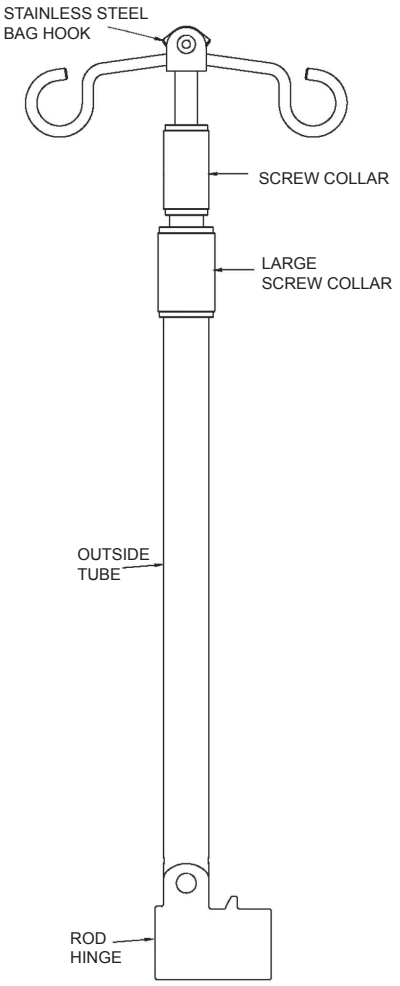
Figure 3.6-1 (STOWED)

Figure 3.6-2 (HINGING)

Figure 3.6-3 (RAISED)

3.7 PERMANENTLY MOUNTED IV ROD OPERATION

Refer to Figure on Right



⚠ **WARNING: PERSONAL INJURY HAZARD - Ensure IV Rod is inserted completely into socket up to the arrow before applying any load.**

3.7.1 Putting IV Rod in UP position

1. Grasp IV Rod and rotate upward until it stops.
2. Push down on IV Rod until it slides firmly into rod hinge socket.

3.7.2 Extending IV Rod

1. Rotate screw collar, or large screw collar, until loosened adequately to allow inner tube to easily slide up or down within outside tube.
2. Lift up on top of IV Rod until desired height is achieved.
3. Tighten collar screw(s) until hand tight.

3.7.3 Retracting IV Rod

1. Support extended portion of IV Rod with one hand.
2. Rotate screw collar until loosened.
3. Lower IV Rod until desired height is achieved, then re-tighten screw collar.
4. Repeat process with second screw collar as required.

3.8 RAIL OPERATION

⚠ **WARNING: Always ensure the rail is locked in position before leaving the patient unattended.**

3.8.1 Raising the Rail

Grasp the rail top cap in the middle of the rail (Figure 3.8-1) and lift.

Figure 3.8-1

3.8.2 Half Height

Grasp the rail and lift the red trigger under the litter top (Figure 3.8-2) while lowering the rail. When the rail starts to move down, release the trigger. Lower the rail until it locks into half height position.

Figure 3.8-2

3.8.3 Lowering the Rail

Grasp the rail and lift the red trigger under the litter top (Figure 3.8.2) while lowering the rail. Continue to lift the trigger until the rail is all the way down.

⚠ **WARNING: When lowering the rails, ensure patient and caregiver body and extremities are clear of pinch points before operating the rail (Figure 3.8-3).**

⚠ **WARNING: Ensure both rails are in upright locked position before leaving patient.**

Figure 3.8-3

4 TROUBLESHOOTING GUIDE

4.1 ELECTRIC POWERED STRETCHERS

⚠ **DANGER: SHOCK HAZARD — To reduce the risk of electric shock, DO NOT remove the cover. Unit is to be serviced by qualified service personnel (minimum 1 year medical equipment service and repair experience) only.**

⚠ **DANGER: SHOCK HAZARD—ALWAYS disconnect the power source whenever troubleshooting or servicing any electric powered stretcher.**

If	Then
One motor or one column does not move but all others are operating correctly.	Step 1: Check all motor and column plug connections at the controller.
	Step 2: If a column does not move: Check the connection at the column.
	Step 3: Plug a connector from the faulty component into a different socket. If the component does not run: Replace that component. If the component runs: Test pendant by plugging a functioning component into the non-functioning socket on the controller. If this component does not run, replace the pendant. If replacing the pendant does not fix the problem, then replace the controller.
Nothing moves.	Step 1: Plug unit into main supply wall receptacle, then observe the pilot light on the controller. If the pilot light is off: Replace the controller. If the pilot light is on: 1. Check the nurse control plug connection at the controller. 2. Check the pendant control plug connection at the controller. 3. Ensure all lockout functions are deactivated. 4. Contact GF Health Products, Inc. for further help and instruction.
The unit runs when plugged into wall receptacle, but does not run on battery backup.	Step 1: Plug unit into a wall receptacle overnight. If the battery doesn't hold a charge, replace the battery (section 4.2).

GF Health Products, Inc. may be contacted at 1.770.368.4700 for additional information required to service or repair the equipment.

4.2 BATTERY REPLACEMENT INSTRUCTIONS

Info: The stretcher utilizes a unique battery specific to this unit. To order, contact Graham-Field Customer Service at 1.770.368.4700.

1. Locate the battery and control box near the front column on the patient right side.
2. Using a flat-head screwdriver, depress the battery mounting tab shown.
3. Remove the control box and battery by sliding them away from you until they become detached.
4. Using a flat-head screwdriver, depress the control box mounting tab.
5. Separate the battery from the control box by pulling toward yourself while supporting the control box.
6. Open the battery cord access cover using a flat-head screwdriver to depress the two locking tabs.
7. Remove the cable from the battery and replace it with a new battery.
8. To reinstall the battery, repeat previous steps in reverse order.

Fig. 4.2-1

Fig. 4.2-2

Fig. 4.2-3

Fig. 4.2-4

Fig. 4.2-5

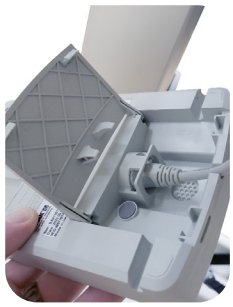


Fig. 4.2-7

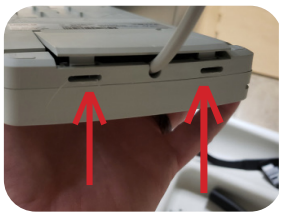


Fig. 4.2-6

5 PREVENTIVE MAINTENANCE FOR THE USER

Component	Cleaning Procedure	Schedule	Cleaning Agent *	Special Notes
Pads / Mattresses	Wipe with damp cloth to remove any foreign material	After each use	Routine hospital grade disinfectants, soap and water	Use only medium strength cleaners Do not steam clean or pressure wash
Stretcher	Wipe with damp cloth to remove any foreign material		Routine hospital grade disinfectants, soap and water	Lubricate pivot points after cleaning
Electrical components	Wipe external surfaces ONLY with damp cloth to remove any foreign material		Routine hospital grade disinfectants, soap and water	Use only medium strength cleaners
Mechanical stretcher components	Wipe with damp cloth to remove any foreign material		Routine hospital grade disinfectants, soap and water	Lubricate pivot points after cleaning
Mechanical accessories	Wipe with damp cloth to remove any foreign material		Routine hospital grade disinfectants, soap and water	Lubricate pivot points after cleaning

Procedure	Schedule	Material
Lubricate all moving and sliding parts and hinge points	Every 3 months	Lubricating oil, light-duty grease, wax stick lubricant or Never-Seez lubricant
⚠ **NEVER LUBRICATE ACTUATORS OR COLUMNS** ⚠		
Inspect all fasteners to ensure proper fit, position and tightness, including nuts, bolts, etc.	Every 3 months	Proper size wrench and screwdriver
Inspect all surfaces and remove any sharp or burred areas; apply touch-up paint where required		Metal file, proper color paint (specify color when ordering)

* Disinfecting and Cleaning Upholstery - ALWAYS follow manufacturer's recommended dilution	
Disinfectants for vinyl products	Phenolic disinfectants are the best choice for vinyl
	Properly diluted quaternaries are also acceptable for vinyl
	Quaternary / Isopropyl disinfectants ARE NOT recommended for vinyl
Disinfectants for urethane products	Quaternary disinfectants are recommended for urethane
	Quaternary / Isopropyl disinfectants are recommended for urethane
	Phenolics SHOULD BE AVOIDED on urethane
Disinfectants for all products	All fabrics may be cleaned with a 1:10 dilution of household bleaches containing 5.25% sodium hypochlorite as recommended by the Centers for Disease Control in Atlanta, Georgia; there is no harmful effect on the fabric
	Disinfectants applied at full concentration or in highly concentrated solutions will decrease the useful life of fabric
	Iodophor-type disinfectants used on fabric may result in staining
Soils or Stains	Use neutral soapsuds and lukewarm water; DO NOT use harsh cleansers, solvents or detergents
Hard-To-Clean Spots	Use standard household / vinyl cleansers and a soft bristle brush on troublesome spots or stains; pre-soak heavy, dried-on soil
Laundering	**Laundering Vinyl-laminated, Polyurethane-coated, or Rubber-coated fabric IS NOT recommended; l**aundering may substantially decrease the useful life of the fabric

▲ **NOTICE — POSSIBLE EQUIPMENT DAMAGE HAZARD: Steam cleaning and pressure washing of stretcher is not recommended and can void warranty.**

Info: *For more detailed information, please contact GF Health Products, Inc. at 1.770.368.4700.*

Info: *GF Health Products, Inc. offers customized Preventive Maintenance Service Programs for Hausted products; contact your GF sales representative for further information.*

6 OPTIONAL ACCESSORIES

Universal Accessories	
H000E1700	IV Rod, 42" Fixed Height
H000018	IV Rod, Telescoping Stainless 27" to 50" Height 2-Section with Holder
HSA080018	IV Pole Pendant Holder
H080770	IV and MONITOR ROD, Telescoping Stainless Steel
HSA400700	Surgical Accessories Rail - Pair
HSA080016	Surgical Bar Adapter for Retracto Rail Models - 5E82 & 5B800
HSA080017	Surgical Arm board (HP141210538) with 2" Navy Pad (HP150830167)
HSA080029	Orthopedic Hand Surgery Board
HSA080015	Acc'y, Patient Safety Strap W/ Buckle and Clip
HSA080021	Next-Gen Oxyflex Diffusion & Extraction System
HSA080041	Dual-Hand Pendant Kit – Stretchers
HSA080010	Mounting Bracket Vertical O2 Tank Holder
H12845000	Vertical O2 Tank Holder
HSA080039	Vertical O2 Tank Holder w/ Mounting Bracket
HSA080014	Acc'y, FS3 Switch - Hi / Lo
HSA080022	Patient Tray - Next-Gen Stretchers
H000N4500	Folding Monitor Shelf
H00WN4500	Folding Monitor Shelf - Extra Wide
HSA080023	Lateral X-Ray Cassette Holder
H06884600	Extension Footboard / Monitor Shelf with Chart Holder – Wide Width
H0CRW6B00	Stationary Footboard / Headboard with Chart Holder – Wide Width
H000W6B00	Stationary Footboard / Headboard – Wide Width
HPD0612	Pad Set, 6 Leg Retracto Rail - Navy
HSA400600	Oxyflex II with Flexible Support Structure and Adapter
HSA007900	Oxyflex II with Flexible Support Structure and Tuck Plate
HSA008000	Disposable Oxyflex II Diffusion Tray Including 24" Tube (Qty 25)

⚠ **WARNING: It is recommended that only accessories approved by GF Health Products, Inc. be used with this device. The use of accessories, transducers, and cables other than those specified by the manufacturer may result in increased emissions or decreased immunity of the Hausted equipment.**

Info: *To order accessories, or for more detailed information on accessories, please contact GF Health Products, Inc. at 1.770.368.4700.*

7 GF HEALTH PRODUCTS, INC. LIMITED WARRANTY FOR HAUSTED

SCOPE OF WARRANTY

GF Health Products, Inc. ("GF") warrants to the original purchaser only that it will replace or repair components, at GF's sole discretion, that are defective in material or workmanship under normal use and service. All warranties are conditioned upon the proper use of the products strictly in accordance with good commercial practice and applicable GF instructions and manuals, including proper use and maintenance. To the extent that a third party warrants a component, GF conveys all of its rights under that warranty to the original purchaser, to the extent permitted.

This limited warranty shall only apply to defects that are reported in accordance with the provisions set forth in this warranty document, within the applicable warranty period and which, upon examination by GF or its authorized representative, prove to be a warranty item. (See Obtaining Warranty Service below) This limited warranty is not transferable.

The warranted components and time periods are set forth below:

COMPONENT	PARTS WARRANTY
Frame	5 years
Casters	1 year
Electrical components	2 years
Hydraulics	1 year
Mechanical Components:	3 years
Original and Replacement Upholstered Tops ǂ:	1 year
Replacement Parts ǂ:	90 days

* Labor is not included in the warranty.

ǂ Upholstery is only warranted on material supplied by GF.

ǂ The warranty period is as designated above. If a part is replaced under warranty, the original warranty period will not be affected. All other replacement parts will be subject to the warranty period specified.

The applicable warranty period shall commence from date of shipment to the original customer, unless there is an expiration date on the component in which case the warranty shall expire on the earlier of warranty period or the expiration date.

OBTAINING WARRANTY SERVICE

Customers located in the United States who wish to report a warranty issue, must contact GF directly by calling 1.770.368.4700 or by e-mailing a request to cs@grahamfield.com. Customers located outside the United States must contact the Distributor from whom they purchased the products. In both cases, further directions will be provided once the initial contact is made. This limited warranty shall only apply to defects that are reported within the applicable warranty period. Failure to abide by the specific directions will result in denial of the warranty claim.

The warranty does not cover and GF shall not be liable for the following:

1) Defects, damage, or other conditions caused, in whole or in part, by misuse, abuse, negligence, alteration, accident, freight damage, tampering or failure to seek and obtain repair or replacement in a timely manner;
2) Products which are not installed, used, or properly cleaned and maintained as required in the official manual for the applicable product;
3) Products considered to be of a non-durable nature including, but not limited to: filters, fuses, gaskets, lubricants, and charts;
4) Accessories or parts not provided by GF;
5) Matching of color, grain or texture except to commercially acceptable standards;
6) Changes in color caused by natural or artificial light:
7) Charges by anyone for adjustments, repairs, replacement parts, installation or other work performed upon or in connection with such products which are not expressly authorized in writing, in advance, by GF;
8) Any labor or shipping charges incurred in the replacement part installation or repair;
9) Costs and expenses of regular maintenance and cleaning; and
10) Representations and warranties made by any person or entity other than GF.

ENTIRE WARRANTY, EXCLUSIVE REMEDY AND CONSEQUENTIAL DAMAGES DISCLAIMER

THIS WARRANTY IS GF'S ONLY WARRANTY AND IS IN LIEU OF ALL OTHER WARRANTIES, EXPRESS OR IMPLIED. GF MAKES NO IMPLIED WARRANTIES OF ANY KIND INCLUDING ANY IMPLIED WARRANTIES OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. IF ANY MODEL OR SAMPLE WAS SHOWN TO THE CUSTOMER, SUCH MODEL OR SAMPLE WAS USED MERELY TO ILLUSTRATE THE GENERAL TYPE AND QUALITY OF THE PRODUCT AND NOT TO REPRESENT THAT THE PRODUCT WOULD NECESSARILY CONFORM TO THE MODEL OR SAMPLE IN ALL RESPECTS. THIS WARRANTY IS LIMITED TO THE REPAIR OR REPLACEMENT OF THE DEFECTIVE PARTS. GF SHALL NOT BE LIABLE FOR AND HEREBY DISCLAIMS ANY DIRECT, SPECIAL, INDIRECT, INCIDENTAL, EXEMPLARY OR CONSEQUENTIAL DAMAGES, INCLUDING, BUT NOT LIMITED TO: DAMAGES FOR LOSS OF PROFITS OR INCOME, LOSS OF USE, DOWNTIME, COVER, OR EMPLOYEE OR INDEPENDENT CONTRACTOR WAGES, PAYMENTS AND BENEFITS. CERTAIN STATES MAY CONFER ADDITIONAL RIGHTS REGARDING WARRANTIES AND IN THOSE STATES GF'S LIABILITY AND THE LIABILITY OF GF'S SUPPLIERS, SHALL BE LIMITED TO THE FULLEST EXTENT PERMITTED BY LAW.

The warranties contained herein, together with GF's current Terms and Conditions, contain all the representations and warranties with respect to the subject matter of this document, and supersede all prior negotiations, agreements and understandings with respect thereto. The recipient of this document hereby acknowledges and represents that it has not relied on any representation, assertion, guarantee, warranty, collateral contract or other assurance, except those set out in this document.

For additional information on this product or this warranty, please contact a GF Customer Service Representative.

NOTES:

1) Additional terms and conditions may apply. See GF's General Terms and Conditions on its website: www.grahamfield.com.
2) Freight claims must be notated on the appropriate shipping documents and must be made with immediacy. International, federal and state regulations govern specific requirements for freight claims. Failure to abide by those regulations may result in a denial of the freight claim. GF will assist you in filing the freight claim.
3) Claims for any short shipment must be made within three (3) days of the invoice date.

8 DISPOSAL AND KEY TO SYMBOLS

DISPOSAL

Hausted equipment and accessories can be disposed of.

We recommend disassembling and dividing the equipment and components into different waste groups such as: metal, cable, electronic, recoverable resource and plastic for recycling or combustion.

Most plastic components are provided with a plastic types code and fiber content to aid sorting of plastic parts.

Product	Metal Scrap	Cable Scrap	Electronic Scrap	Plastic Recycling or Combustion
5B800ST	X	X	X	X

Info: Dispose of lithium battery attached to the control box in accordance with local regulations.

KEY TO SYMBOLS

The following symbols are used on Hausted product labels.

	Protective Earth		Manufacturer
	Earth Ground		Keep Dry
	General Warning Sign		Fragile, Handle with Care
	CE Mark		Electrical and Electronic Equipment
	ETL		Consult Instructions for Use
EU REP	European Authorized Representative		Caution
	Disconnect before Service		Pinch Point
MD	Medical Device	UDI	Unique Device Identifier
	Type B Applied Part		

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9 APPENDIX

9.1 GUIDANCE AND MANUFACTURER'S DECLARATION — ELECTROMAGNETIC EMISSIONS

The Hausted Stretchers are intended for use in the electromagnetic environment specified below. The customer or the user of the Hausted Stretchers should assure that they are used in such an environment.

Emissions Test	Compliance	Electromagnetic Environment — Guidance
RF emissions CISPR 11	Group 1	The Hausted Stretchers use RF energy only for their internal function. Therefore, their RF emissions are very low and are not likely to cause any interference in nearby electronic equipment. The Hausted Stretchers are suitable for use in all establishments other than domestic and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
RF emissions CISPR 11	Class A	
Harmonic emissions IEC 61000-3-2	Class A	
Voltage fluctuations / flicker emissions IEC 61000-3-3	Complies	

9.2 ENCLOSURE PORT [1]

Phenomenon	Basic EMC standard or test method	IMMUNITY TEST LEVELS
		Professional healthcare facility environment
ELECTROSTATIC DISCHARGE	IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air
Radiated RF EM fields [a]	IEC 61000-4-3	3 V/m [f] 80 MHz – 2,7 GHz [b] 80 % AM at 1 kHz [c]
Proximity fields from RF wireless communications equipment	IEC 61000-4-3	See Table 9.3.
RATED power frequency magnetic fields [d] [e]	IEC 61000-4-8	30 A/m [g] 50 Hz or 60 Hz

[a] The interface between the PATIENT physiological signal simulation, if used, and the ME EQUIPMENT or ME SYSTEM shall be located within 0,1 m of the vertical plane of the uniform field area in one orientation of the ME EQUIPMENT or ME SYSTEM.

[b] ME EQUIPMENT and ME SYSTEMS that intentionally receive RF electromagnetic energy for the purpose of their operation shall be tested at the frequency of reception. Testing may be performed at other modulation frequencies identified by the RISK MANAGEMENT PROCESS. This test assesses the BASIC SAFETY and ESSENTIAL PERFORMANCE of an intentional receiver when an ambient signal is in the passband. It is understood that the receiver might not achieve normal reception during the test.

[c] Testing may be performed at other modulation frequencies identified by the RISK MANAGEMENT PROCESS.

[d] Applies only to ME EQUIPMENT and ME SYSTEMS with magnetically sensitive components or circuitry.

[e] During the test, the ME EQUIPMENT or ME SYSTEM may be powered at any NOMINAL input voltage, but with the same frequency as the test signal (see Table 1).

[f] Before modulation is applied.

[g] This test level assumes a minimum distance between the ME EQUIPMENT or ME SYSTEM and sources of power frequency magnetic field of at least 15 cm. If the RISK ANALYSIS shows that the ME EQUIPMENT or ME SYSTEM will be used closer than 15 cm to sources of power frequency magnetic field, the IMMUNITY TEST LEVEL shall be adjusted as appropriate for the minimum expected distance.

9.3 ENCLOSURE PORT IMMUNITY TO RF WIRELESS COMMUNICATIONS EQUIPMENT [1]

Test frequency (MHz)	Band [a] (MHz)	Service [a]	Modulation [b]	Maximum power (W)	Distance (m)	IMMUNITY TEST LEVEL (V/m)
385	380 –390	TETRA 400	Pulse modulation [b] 18 Hz	1,8	0,3	27
450	430 – 470	GMRS 460, FRS 460	FM [c] ± 5 kHz deviation 1 kHz sine	2	0,3	28
710	704 – 787	LTE Band 13, 17	Pulse modulation [b] 217 Hz	0,2	0,3	9
745						
780						
810	800 – 960	GSM 800/900, TETRA 800, iDEN 820, CDMA 850, LTE Band 5	Pulse modulation [b] 18 Hz	2	0,3	28
870						
930						
1 720	1 700 – 1 990	GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3, 4, 25; UMTS	Pulse modulation [b] 217 Hz	2	0,3	28
1 845						
1 970						
2 450	2 400 – 2 570	Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse modulation [b] 217 Hz	2	0,3	28
5 240	5 100 – 5 800	WLAN 802.11 a/n	Pulse modulation [b] 217 Hz	0,2	0,3	9
5 500						
5 785						

9.4 INPUT AC POWER PORT [1]

Phenomenon	Basic EMC standard	IMMUNITY TEST LEVELS
		Professional healthcare facility environment
Electrical fast transients / bursts [a) l) o)]	IEC 61000-4-4	± 2 kV 100 kHz repetition frequency
Surges [a) b) j) o)] Line-to-line	IEC 61000-4-5	± 0,5 kV, ± 1 kV
Surges [a) b) j) k) o)] Line-to-ground	IEC 61000-4-5	± 0,5 kV, ± 1 kV, ± 2 kV
Conducted disturbances induced by RF fields [c) d) o)]	IEC 61000-4-6	3 V [m)] 0,15 MHz – 80 MHz 6 V [m)] in ISM bands between 0,15 MHz and 80 MHz [n)] 80 % AM at 1 kHz [e)]
Voltage dips [f) p) r)]	IEC 61000-4-11	0 % U_T; 0,5 cycle [g)] At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° [q)]
		0 % U_T; 1 cycle and 70 % U_T; 25/30 cycles [h)] Single phase: at 0°
Voltage interruptions [f) i) o) r)]	IEC 61000-4-11	0 % U_T; 250/300 cycle [h)]

a) The test may be performed at any one power input voltage within the ME EQUIPMENT or ME SYSTEM RATED voltage range. If the ME EQUIPMENT or ME SYSTEM is tested at one power input voltage, it is not necessary to re-test at additional voltages.

b) All ME EQUIPMENT and ME SYSTEM cables are attached during the test.

c) Calibration for current injection clamps shall be performed in a 150 Ω system.

d) If the frequency stepping skips over an ISM or amateur band, as applicable, an additional test frequency shall be used in the ISM or amateur radio band. This applies to each ISM and amateur radio band within the specified frequency range.

e) Testing may be performed at other modulation frequencies identified by the RISK MANAGEMENT PROCESS.

f) ME EQUIPMENT and ME SYSTEMS with a d.c. power input intended for use with a.c.-to-d.c. converters shall be tested using a converter that meets the specifications of the MANUFACTURER of the ME EQUIPMENT or ME SYSTEM. The IMMUNITY TEST LEVELS are applied to the a.c. power input of the converter.

g) Applicable only to ME EQUIPMENT and ME SYSTEMS connected to single-phase a.c. mains.

h) E.g. 10/12 means 10 periods at 50 Hz or 12 periods at 60 Hz.

i) ME EQUIPMENT and ME SYSTEMS with RATED input current greater than 16 A / phase shall be interrupted once for 250/300 cycles at any angle and at all phases at the same time (if applicable). ME EQUIPMENT and ME SYSTEMS with battery backup shall resume line power operation after the test. For ME EQUIPMENT and ME SYSTEMS with RATED input current not exceeding 16 A, all phases shall be interrupted simultaneously.

9.4 CONTINUED

j) ME EQUIPMENT and ME SYSTEMS that do not have a surge protection device in the primary power circuit may be tested only at ± 2 kV line(s) to earth and ± 1 kV line(s) to line(s).

k) Not applicable to CLASS II ME EQUIPMENT and ME SYSTEMS.

l) Direct coupling shall be used.

m) r.m.s., before modulation is applied.

n) The ISM (industrial, scientific and medical) bands between 0,15 MHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; and 40,66 MHz to 40,70 MHz. The amateur radio bands between 0,15 MHz and 80 MHz are 1,8 MHz to 2,0 MHz, 3,5 MHz to 4,0 MHz, 5,3 MHz to 5,4 MHz, 7 MHz to 7,3 MHz, 10,1 MHz to 10,15 MHz, 14 MHz to 14,2 MHz, 18,07 MHz to 18,17 MHz, 21,0 MHz to 21,4 MHz, 24,89 MHz to 24,99 MHz, 28,0 MHz to 29,7 MHz and 50,0 MHz to 54,0 MHz.

o) Applicable to ME EQUIPMENT and ME SYSTEMS with RATED input current less than or equal to 16 A / phase and ME EQUIPMENT and ME SYSTEMS with RATED input current greater than 16 A / phase.

p) Applicable to ME EQUIPMENT and ME SYSTEMS with RATED input current less than or equal to 16 A / phase.

q) At some phase angles, applying this test to ME EQUIPMENT with transformer mains power input might cause an overcurrent protection device to open. This can occur due to magnetic flux saturation of the transformer core after the voltage dip. If this occurs, the ME EQUIPMENT or ME SYSTEM shall provide BASIC SAFETY during and after the test.

r) For ME EQUIPMENT and ME SYSTEMS that have multiple voltage settings or auto ranging voltage capability, the test shall be performed at the minimum and maximum RATED input voltage. ME EQUIPMENT and ME SYSTEMS with a RATED input voltage range of less than 25 % of the highest RATED input voltage shall be tested at one RATED input voltage within the range. See Table 1 Note c) for examples calculations.

9.5 PATIENT COUPLING PORT [1]

Phenomenon	Basic EMC standard	IMMUNITY TEST LEVELS
		Professional healthcare facility environment
ELECTROSTATIC DISCHARGE [c)]	IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air
Conducted disturbances induced by RF fields [a)]	IEC 61000-4-6	3 V [b)] 0,15 MHz – 80 MHz 6 V [b)] in ISM bands between 0,15 MHz and 80 MHz 80 % AM at 1 kHz

a) The following apply:

- All PATIENT-COUPLED cables shall be tested, either individually or bundled
- PATIENT-COUPLED cables shall be tested using a current clamp unless a current clamp is not suitable. In cases were a current clamp is not suitable, an EM clamp shall be used.
- No intentional decoupling device shall be used between the injection point and the PATIENT COUPLING POINT in any case.
- Testing may be performed at other modulation frequencies identified by the RISK MANAGEMENT PROCESS.
- Tubes that are intentionally filled with conductive liquids and intended to be connected to a PATIENT shall be considered to be PATIENT-COUPLED cables.
- If the frequency stepping skips over an ISM or amateur radio band, as applicable, an additional test frequency shall be used in the ISM or amateur radio band. This applies to each ISM and amateur radio band within the specified frequency range.
- The ISM (industrial, scientific and medical) bands between 0,15 MHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; and 40,66 MHz to 40,70 MHz. The amateur radio bands between 0,15 MHz and 80 MHz are 1,8 MHz to 2,0 MHz, 3,5 MHz to 4,0 MHz, 5,3 MHz to 5,4 MHz, 7 MHz to 7,3 MHz, 10,1 MHz to 10,15 MHz, 14 MHz to 14,2 MHz, 18,07 MHz to 18,17 MHz, 21,0 MHz to 21,4 MHz, 24,89 MHz to 24,99 MHz, 28,0 MHz to 29,7 MHz and 50,0 MHz to 54,0 MHz.

b) r.m.s., before modulation is applied

c) Discharges shall be applied with no connection to an artificial hand and no connection to PATIENT simulation. PATIENT simulation may be connected after the test as needed in order to verify BASIC SAFETY and ESSENTIAL PERFORMANCE.

9.6 SIGNAL INPUT/OUTPUT PARTS PORT [1]

Phenomenon	Basic EMC standard	IMMUNITY TEST LEVELS
		Professional healthcare facility environment
ELECTROSTATIC DISCHARGE [e]	IEC 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air
Electrical fast transients / bursts [b] [f]	IEC 61000-4-4	± 1 kV 100 kHz repetition frequency
Surges Line-to-ground [a]	IEC 61000-4-5	± 2 kV
Conducted disturbances induced by RF fields [b] [d] [g]	IEC 61000-4-6	3 V [h] 0,15 MHz – 80 MHz 6 V [h] in ISM bands between 0,15 MHz and 80 MHz [i] 80 % AM at 1 kHz [c]

[a] This test applies only to output lines intended to connect directly to outdoor cables.

[b] SIP/SOPS whose maximum cable length is less than 3 m in length are excluded.

[c] Testing may be performed at other modulation frequencies identified by the RISK MANAGEMENT PROCESS.

[d] Calibration for current injection clamps shall be performed in a 150 Ω system.

[e] Connectors shall be tested per 8.3.2 and Table 4 of IEC 61000-4-2:2008. For insulated connector shells, perform air discharge testing to the connector shell and the pins using the rounded tip finger of the ESD generator, with the exception that the only connector pins that are tested are those that can be contacted or touched, under conditions of INTENDED USE, by the standard test finger shown in Figure 6 of the general standard, applied in a bent or straight position.

[f] Capacitive coupling shall be used.

[g] If the frequency stepping skips over an ISM or amateur radio band, as applicable, an additional test frequency shall be used in the ISM or amateur radio band. This applies to each ISM and amateur radio band within the specified frequency range.

[h] r.m.s., before modulation is applied.

[i] The ISM (industrial, scientific and medical) bands between 150 kHz and 80 MHz are 6,765 MHz to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; and 40,66 MHz to 40,70 MHz. The amateur radio bands between 0,15 MHz and 80 MHz are 1,8 MHz to 2,0 MHz, 3,5 MHz to 4,0 MHz, 5,3 MHz to 5,4 MHz, 7 MHz to 7,3 MHz, 10,1 MHz to 10,15 MHz, 14 MHz to 14,2 MHz, 18,07 MHz to 18,17 MHz, 21,0 MHz to 21,4 MHz, 24,89 MHz to 24,99 MHz, 28,0 MHz to 29,7 MHz and 50,0 MHz to 54,0 MHz.

5B800ST-PD NAVIGATOR SMART DRIVE SYSTEM ADDENDUM

CONTENTS

1 PRODUCT DESCRIPTION

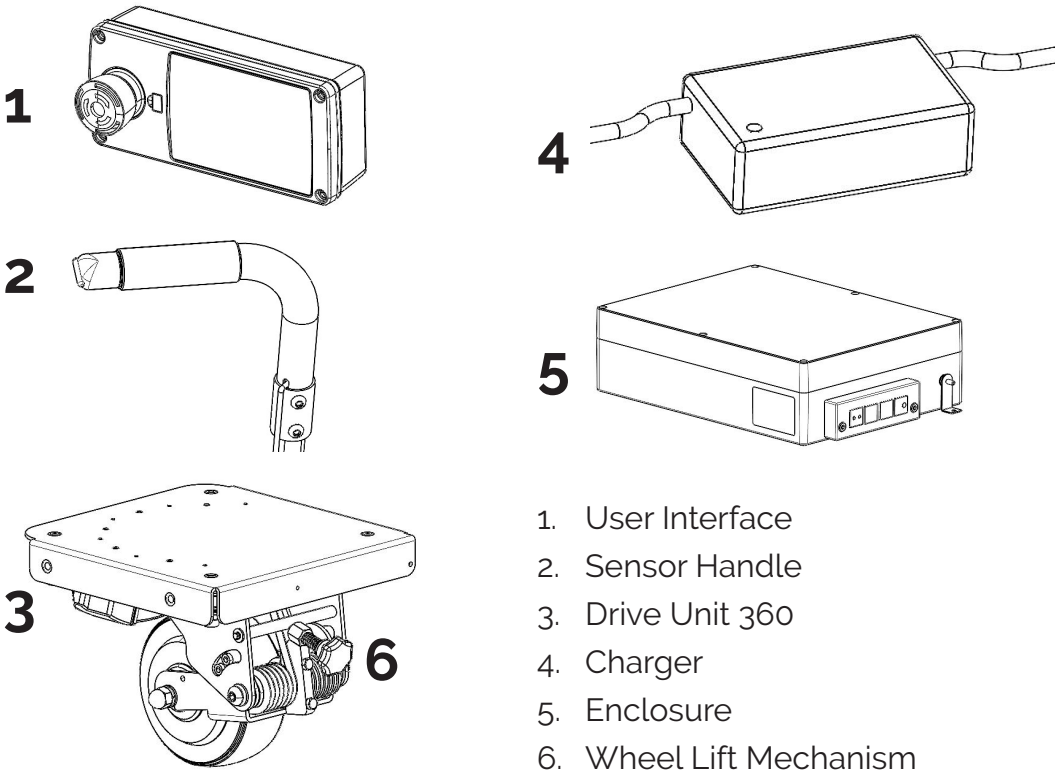
This manual describes use of the Navigator Smart Drive System (“NSD”) power assist. NSD is an add-on solution for stretcher products.

The system features an electric drive wheel that provides power assist drive functionality for forwards, backwards and sideways movement. Power assist means that the user uses some force to push and pull the stretcher. The system multiplies these forces to drive the wheel. With this technique, the NSD makes it possible to move stretchers in a natural way, whether the routes are long, on carpet, over thresholds, on slopes or while maneuvering in small spaces. Heavy stretchers are moved around as if they only weigh a couple of pounds.

▲ **NOTICE: The NSD only concerns the functionality for electric power assist drive of the stretcher. Please refer to the user manual of the stretcher for any information regarding its complete functionality.**

2 COMPONENT OVERVIEW

Each system consists of a set of components. These components may be visible or hidden, depending on how they are built on the stretcher. Below table lists components that may be identified on your stretcher.



1. User Interface
2. Sensor Handle
3. Drive Unit 360
4. Charger
5. Enclosure
6. Wheel Lift Mechanism

In most cases, the Sensor Handle and User Interface are the only components the user directly interacts with. Please contact your supplier for further details of integration of NSD components on your stretcher.

3 LIST OF WARNINGS AND CAUTIONS

⚠ **IMPORTANT: Read this manual completely before use of stretchers equipped with the NSD.**

⚠ **CAUTION: Stretchers equipped with the NSD may only be used by professional, well trained and adult users aware of all instructions in this manual.**

⚠ **CAUTION: When using the NSD, at all times stay focused on the task of maneuvering the stretcher. Do not use the NSD when focus on this task is not possible for any reason or when using any medication for which it is advised not to use machines or participate in traffic.**

⚠ **CAUTION: Make sure the intended route of the stretcher is free of obstacles and people.**

⚠ **CAUTION: Always ensure good sight in walking direction during movement of heavy stretchers.**

⚠ **CAUTION: Adjust the speed of the stretcher to a level that makes it possible to respond timely on (suddenly) oncoming people and obstacles.**

⚠ **WARNING: Do not use stretchers with NSD on slopes exceeding a 6 degree angle. Stretchers with NSD must not be used on slopes unsuitable for unpowered pushing and pulling of stretchers.**

⚠ **WARNING: The NSD ensures that heavy stretchers can be maneuvered with much lower push and pull forces. Please make sure to adjust your driving style. Activation of the system only requires minimal forces on the sensor handle thumb switch. Avoid high and sudden push and pull forces on the sensor handle.**

⚠ **CAUTION: Always switch the NSD off when the stretcher is not in use or is left unattended.**

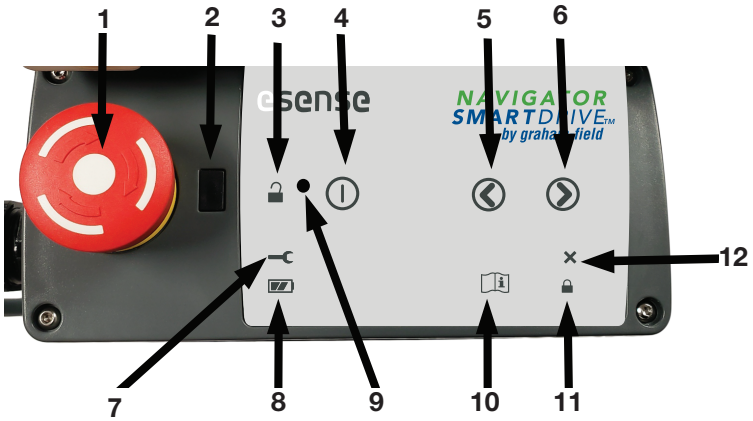
⚠ **CAUTION: Make sure the NSD is not used in combination with other electrical or mechanical means for propulsion of the stretcher. These may harm the NSD components.**

⚠ **CAUTION: The NSD is controlled by a sensitive sensor in the right sensor handle. Ensure at all times that the sensor handle thumb switch can move freely and no forces are exerted on it other than the forces required for the continued pushing of the stretcher.**

- **Prevent objects from pushing against the sensor handle thumb switch.**
- **Do not hang clothes, bags, etc. on the handles. Take care that no objects are clamped between the sensor handle thumb switch and other surfaces.**

4 NAVIGATOR SMART DRIVE ("NSD") USE

4.1 USER INTERFACE ELEMENTS



Elements in table (see right) can be identified.

1. Emergency Switch	Active
2. USB Connector	Active
3. Magnet Key Lock	Not Active
4. On / Off Switch	Active
5. Move Left	Active
6. Move Right	Active
7. Service Indication (But Only Lit In Case of Service Errors)	Active,
8. Battery Indication	Active
9. System Status Indication	Active
10. Consult Manual	Reference
11. Key Lock Indication	Not Active
12. Immobilizer Indication	Active

4.2 PREPARATIONS

The NSD system is activated by pushing the On / Off switch briefly. At start-up a short audible beep will sound. Directly after activation, the system requires a few seconds to calibrate the force sensor inside the sensor handle. Additionally, the direction of the Drive wheel is automatically checked by the system.

During calibration the system status indication blinks green. When calibration is finished a second audible beep will sound and the system status indication changes to continuously lit green. The system is now ready for use.

⚠ **CAUTION: Before switching the system on, make sure that the sensor handle thumb switch is completely free of external forces.**

- **Prevent objects, such as blankets, to push against the sensor handle thumb switch.**
- **Do not hang clothes, bags, etc. on the handles.**

  In extreme situations such external forces during calibration may lead to unexpected behavior of the drive system.

▲ **NOTICE: Do not place your hand on the sensor handle immediately after activation of the NSD. The sensor handle thumb switch shall not be moved or touched during calibration. If this happens the system will postpone calibration until the sensor handle is released. When calibration is postponed by more than ten seconds a warning beep sounds every two seconds. Remove hand or other objects from the sensor handle in this situation. The system status indication will continue to blink green until calibration is performed successfully.**
In case calibration was not successful within one minute the system will switch to service error mode. See paragraph 'Service mode' for reference. The drive system is switched off. Failure of calibration may be caused by an external reason, as described above, but may also be caused by a failure in the sensor handle thumb switch. Please contact your stretcher supplier in case the error persists.

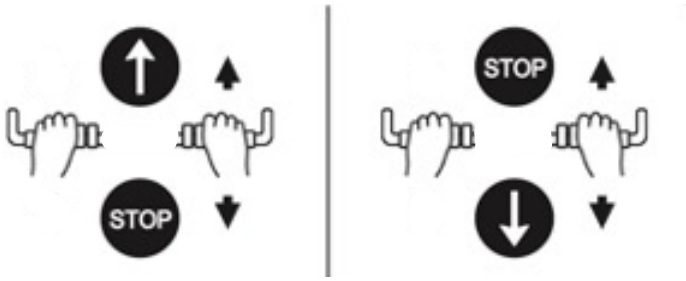
▲ **NOTICE: Make sure the brakes of the stretcher are released, and the casters are in the neutral position before use of the system.**

▲ **NOTICE: The steer mode function (by pendant or manually activated) is disabled on stretchers equipped with the Navigator Smart Drive System. See page 14.**

▲ **NOTICE: Make sure the emergency switch is off before using the unit for the first time. The emergency switch is locked in the on position during shipment. The emergency switch can be unlocked by first turning it clockwise and secondly pressing the on / off switch briefly.**

4.3 DRIVING

Power assisted forwards and backwards driving of the stretcher is done in the same manner as forwards and backwards driving of a stretcher without an NSD system. The power assisted drive is automatically activated by placing the right-hand thumb on the thumb control sensor located on the Head-End Patient Right Push Handle. By applying thumb pressure up and down, it will move the stretcher forward and backward. The speed is proportional to the amount of pressure applied.



Power assisted driving is by standard maximized on a speed of 3.62 km / h (2.25 mph).

▲ **NOTICE: The stretcher can be braked actively by practicing force on the thumb control sensor in opposite direction of the direction of travel.**

The NSD system also provides powered assist drive when moving a stretcher up and down on slopes. In case a stretcher is released on a slope it will brake automatically and will gradually drive down slowly.

⚠ **WARNING: Never use stretchers with NSD on slopes exceeding a 6 degree angle.**

▲ **NOTICE: Always apply gentle pressure to the thumb control sensor. In exceptional situations, if too much force is put on the sensor, the system automatically switches to service error mode and drive functionality is switched off. In this case the power assisted drive is switched off and service indication lights are lit. To reset the system in this situation the thumb control sensor shall be released, and the system shall be switched off and on again.**

4.4 SIDEWAYS DRIVING

The stretcher can be maneuvered 90 degrees sideways with power assisted drive.

User position during forwards and backwards driving is the same for sideways driving.

Push and hold the left arrow button to move the stretcher to the left at a constant slow speed and push & hold the right arrow button to move the stretcher to the right at a constant slow speed.

▲ **NOTICE: In case the stretcher is still driving forwards or backwards, the sideways driving functionality on NSD systems will not work immediately. Pushing the left & right arrow buttons for sideways driving will first actively brake the stretcher to a standstill before the stretcher starts driving sideways.**

4.5 EMERGENCIES

The NSD system is equipped with an emergency switch. Pressing this switch immediately switches off power to the NSD system. The switch shall be used only when control over the stretcher is lost.

After pressing the emergency switch all indication lights will start blinking. The emergency switch can be unlocked by turning it clockwise. After unlocking, all indication lights will remain blinking. The system can be reactivated by pressing On / Off switch briefly.

⚠ **WARNING: The emergency switch is not a brake. The switch is intended to switch the drive wheel off. Any power assist drive or braking is immediately stopped and the drive wheel switches to free wheel mode.**

Special notice shall be taken that this can lead to a sudden high increase in required push and pull forces when driving up and down slopes. Any active braking on slopes will also be stopped, causing the stretcher to roll on through.

▲ **NOTICE: The emergency switch is subject to wear and is not designed for switching off the system. Only use the switch in case of emergency. Use the on/ off switch to switch the system off.**

4.6 BATTERY LEVEL INDICATION

The NSD system is provided with an intelligent BMS (Battery Management System) battery that provides sufficient capacity for over 15km (9 miles) of power assist drive. For these systems, the battery indication on the user-interface shows the approximate percentage of battery capacity left.

For this system, the battery indication consists of 5 lights; one orange, one yellow and three green. Below table shows the meaning of the various indications.

Battery indication		Capacity left	Remarks
🔋 🟠 🟡 🟢 🟢 🟢		81 – 100 %	
🔋 🟠 🟡 🟢 🟢 ○		61 – 80 %	
🔋 🟠 🟡 🟢 ○ ○		41 – 60 %	
🔋 🟠 🟡 ○ ○ ○		21 – 40 %	
🔋 🟠 ○ ○ ○ ○		11 – 20 %	Charging advised
🔋 🟠 ○ ○ ○ ○ ♫	*	0 – 10 %	Immediate charging required

* Orange light blinking and beep sounds every 20 seconds.

Battery indication is displayed only when the system is switched on.

4.7 CHARGING

The NSD system is provided with a charger. This charger is integrated on the stretcher. The battery will automatically charge when the stretcher is connected to the mains socket.

The battery provides power for up to 9 miles of driving in most use situations. It's advised to recharge the battery on a regular basis or when the battery indication is lit continuously orange.

▲ **NOTICE: In case the battery indication starts blinking orange and a beep sounds every 20 seconds the battery is almost empty and immediate charging is required.**

Charging of the battery can both be done with the NSD system switched off and when the system is switched on. When the system is switched on during charging, a charge indication is shown on the user-interface and drive functionality will be switched off. In case the thumb control sensor is pressed during charging a beep will sound every two seconds.

It can take up to five seconds before the system notes the charger status and starts or stops charge indication and switches drive functionality off or on.

The charge indication is an animation with increasing number of battery indication lights. The number of lights that remains continuously lit indicates the charge level of the battery.

Charge indication	Description	Charge level
	All 5 lights animate	0 – 10 %
	First light continuously lit, other lights animate	20 – 39 %
	First two lights continuously lit, other lights animate	40 – 59 %
	First three lights continuously lit, other lights animate	60 – 79 %
	Only last light blinks	80 – 100 %

When the battery is fully charged, the fifth battery indication light remains blinking on double speed. Drive functionality remains switched off as long as the stretcher or charger is connected to the mains.

Charging of a fully empty battery takes up to 12 hours. It is advised to charge the battery overnight on a regular basis.

4.8 IMMOBILIZER

The immobilizer function offers a simple way to prevent activation of the power assist drive when the stretcher is not ready for use. For the NSD system, three conditions must be met:

- **The brakes are unlocked.**
- **The stretcher is level (both columns the same height).**
- **The power cable must be unplugged from the wall outlet.**

When the immobilizer is switched on the immobilizer indication will be lit. Power assist drive is switched off and a beep will sound every two seconds when the thumb control sensor is pressed.

4.9 TIMERS

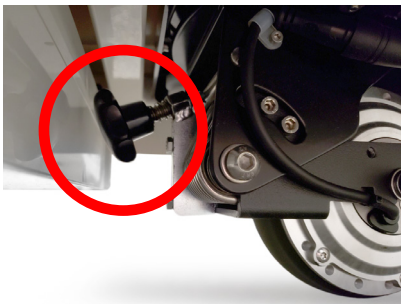
ALL NSD systems are provided with an automatic switch off function. The system will go into energy-save mode at one minute and at two minutes of inactivity the system will automatically switch off. A full restart is required to continue driving.

4.10 AFTER USE

Always switch the NSD system off after use or when the stretcher is left unattended. The system can be switched off by pressing the On / Off switch for a second. All lights on the user-interface will turn off.

4.11 WHEEL LIFT MECHANISM

In the event of a Power Drive Failure, the stretcher will perform normally when manually pushing forwards & backwards. Side-to-Side movement will not be possible. For ease of side-to-side movement, the NSD system is equipped with a wheel lift mechanism that will allow the power assist wheel to be raised from the floor by turning the knob in a clockwise direction until the wheel is off the floor.



5 MAINTENANCE

The thumb control sensor, user-interface, and power assist wheel can be cleaned with a well-wrung wet cloth. Use an ordinary, non-abrasive detergent (PH 5.5-8).

Regularly check the axis and tread of the power assist wheel for accumulation of hair and dust. Carefully remove any debris when needed.

⚠ **CAUTION: Regularly check the NSD system components for loose parts and wear. In case of any irregularities stop use of the system and contact your supplier.**

⚠ **CAUTION: Always switch the system off before maintenance is carried out.**

⚠ **CAUTION: Maintenance on the electrical system shall be performed by qualified service personnel only.**

▲ **NOTICE: Avoid the use of aggressive detergents / solvents and excess of water. The NSD system components are not designed for exposure to large amounts (jets) of water. The NSD system is not suitable for machine washing.**

▲ **NOTICE: Even when the NSD system is not being used, the battery will slowly drain. A fully drained battery can get damaged. Such damage is excluded from the guarantee. It is strongly recommended to fully charge the battery at least every three months. Damage to the battery due to not charging for long periods of time can be monitored and is excluded from warranty.**

6 SERVICE MODE

If the NSD system encounters an error during use that cannot be automatically repaired, it switches to service mode. Power assist drive is switched off, the system status indication on the user-interface will blink orange and one or more service indication lights will be lit.

The pattern of service lights represents a code for the cause of the error.

Service indication	Number	Description
	1	System error
	2	Push Handle error
	3	Interface error
	4	Drive error
	5	Steer error
	6	Battery error

In case your system switches to service mode, first switch the system off and on again. Please contact your supplier in case the error persists or in case the error is noted frequently.

Your supplier may ask for the code of the service indication. Please use below table for reference.

7 TROUBLESHOOTING

In case your NSD system fails operation, perform the following steps:

1. Check if the system is still switched on. The system switches off automatically when it is not used for a period of time. This period is set on two minutes by default.
2. See section 4.8 immobilizer.
3. Make sure the thumb control sensor is not touched for a few seconds directly after starting the system. The sensor needs some time for calibration. In case the thumb control sensor is touched during calibration the calibration will be postponed. If calibration is postponed for more than ten seconds an audible beep will sound every two seconds. If calibration fails for one minute the system will switch to service mode and a full restart is required.
4. Switch the system off and on again. Let the system reboot and check if the system operates normal again.
5. Check if the emergency switch is pressed. All lights on the user interface will blink if this is the case. The emergency switch can be unlocked by first turning it clockwise and secondly pressing the on / off switch briefly.
6. Check the remaining battery power. If the orange light on the battery indication is blinking charging of the battery is required.
7. Check if the stretcher is connected to the mains with a power cord. During charging the battery indication will animate as described in paragraph 'Charging' and power assist drive is switched off. Disconnect the power cord and check if the system operates normal again.
8. Check if one or more service indication lights remain lit after switching the system off and on again. If this is the case the system is switched to service mode and power assist drive is switched off. Please contact your supplier in case the error persists or in case the error is noted frequently.

Your supplier may ask for the code of the service indication. Please use the table in paragraph 'Service mode' for reference.

In case the NSD system operates unexpectedly, perform the following steps:

1. Switch the NSD system off and on again. Let the system restart and then check if the system operates normally. Please contact your supplier in case the problem persists.
2. Check if the thumb control sensor can be pressed freely and make sure no objects are connected to or hanging from the push handles. Remove any obstacles and check if the system resumes normal operation.

Please contact your supplier in case above steps did not solve the problem.

8 PRODUCT SPECIFICATION

Use Environment	
• Duty Cycle 10% 9 min on / 81 min off	The NSD is suited for indoor use only on hard (solid), dry, and clean industrial surfaces.

Ambient Use Temperature	
• Charging	10° to 40° C
• Discharging (Power Assisted Driving)	-20° to 50° C

Storage Temperature	
• Longer Period	-20° to 35° C
• Up to One Week	-20° to 60° C

Charge Time for Full-Charge	12 h
Action Radius on Full Battery	> 15 km
Drive Wheel Functionality When Powered Off	Freewheel, No Parking Brake
Max. Assisted Speed	Up to 3.62 km / h (2.25 mph)
Max. Wheel Torque Continuous	15 Nm
Max. Wheel Torque Short	>30 Nm
Max. Slope Angle (At stretcher weight up to 400 kg)	6°
Max. Threshold	20 mm
Average Braking Distance at Release of Stretcher Sensor Handle*	1.0 m (39 in.)
Average Braking Distance for Active Braking*	0.8 m

Protection Rating of External Components	
• Sensor Handle	IPx5
• User Interface	IPx4
• Drive Unit 360	IPx4
• Controller Box	IPx4

Average Power Consumption	
• Standby	< 0.5 W
• Rest	< 2.5 W
• Horizontal Plane / Acceleration*	72 W
• Horizontal Plane / Constant Speed	24 W
• At Max. Torque	480 W / max. 10 seconds 360 W / max. 5 minutes 240 W / Continuous

Battery	
• Over-temperature Cutoff	45°C
• Low Voltage Cutoff	20V
• Over-voltage Cutoff	34V
• Over-current Cutoff	22A
• Charge Over-current Cutoff	3.5A

* All measured at speed accelerating to approximately. 3.62 km/h (2.25 mph). Total weight of stretcher and load 400 kg.

9 GF HEALTH PRODUCTS, INC. LIMITED WARRANTY FOR HAUSTED

SCOPE OF WARRANTY

GF Health Products, Inc. ("GF") warrants to the original purchaser only that it will replace or repair components, at GF's sole discretion, that are defective in material or workmanship under normal use and service. All warranties are conditioned upon the proper use of the products strictly in accordance with good commercial practice and applicable GF instructions and manuals, including proper use and maintenance. To the extent that a third party warrants a component, GF conveys all of its rights under that warranty to the original purchaser, to the extent permitted. This limited warranty shall only apply to defects that are reported in accordance with the provisions set forth in this warranty document, within the applicable warranty period and which, upon examination by GF or its authorized representative, prove to be a warranty item. (See Obtaining Warranty Service below) This limited warranty is not transferable.

The warranted components and time periods are set forth below:

COMPONENT	**PARTS WARRANTY**
Frame	5 years
Casters	1 year
Electrical components	2 years
Hydraulics	1 year
Mechanical Components:	3 years
Original and Replacement Upholstered Tops ǂ:	1 year
Replacement Parts ǂ:	90 days

* Labor is not included in the warranty.

ǂ Upholstery is only warranted on material supplied by GF.

ǂ The warranty period is as designated above. If a part is replaced under warranty, the original warranty period will not be affected. All other replacement parts will be subject to the warranty period specified.

The applicable warranty period shall commence from date of shipment to the original customer, unless there is an expiration date on the component in which case the warranty shall expire on the earlier of warranty period or the expiration date.

OBTAINING WARRANTY SERVICE

Customers located in the United States who wish to report a warranty issue, must contact GF directly by calling 1.770.368.4700 or by e-mailing a request to cs@grahamfield.com. Customers located outside the United States must contact the Distributor from whom they purchased the products. In both cases, further directions will be provided once the initial contact is made. This limited warranty shall only apply to defects that are reported within the applicable warranty period. Failure to abide by the specific directions will result in denial of the warranty claim.

The warranty does not cover and GF shall not be liable for the following:

1. Defects, damage, or other conditions caused, in whole or in part, by misuse, abuse, negligence, alteration, accident, freight damage, tampering or failure to seek and obtain repair or replacement in a timely manner;
2. Products which are not installed, used, or properly cleaned and maintained as required in the official manual for the applicable product;
3. Products considered to be of a non-durable nature including, but not limited to: filters, fuses, gaskets, lubricants, and charts;
4. Accessories or parts not provided by GF;
5. Matching of color, grain or texture except to commercially acceptable standards;
6. Changes in color caused by natural or artificial light:
7. Charges by anyone for adjustments, repairs, replacement parts, installation or other work performed upon or in connection with such products which are not expressly authorized in writing, in advance, by GF;
8. Any labor or shipping charges incurred in the replacement part installation or repair;
9. Costs and expenses of regular maintenance and cleaning; and
10. Representations and warranties made by any person or entity other than GF.

ENTIRE WARRANTY, EXCLUSIVE REMEDY AND CONSEQUENTIAL DAMAGES DISCLAIMER

THIS WARRANTY IS GF'S ONLY WARRANTY AND IS IN LIEU OF ALL OTHER WARRANTIES, EXPRESS OR IMPLIED. GF MAKES NO IMPLIED WARRANTIES OF ANY KIND INCLUDING ANY IMPLIED WARRANTIES OF MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE. IF ANY MODEL OR SAMPLE WAS SHOWN TO THE CUSTOMER, SUCH MODEL OR SAMPLE WAS USED MERELY TO ILLUSTRATE THE GENERAL TYPE AND QUALITY OF THE PRODUCT AND NOT TO REPRESENT THAT THE PRODUCT WOULD NECESSARILY CONFORM TO THE MODEL OR SAMPLE IN ALL RESPECTS. THIS WARRANTY IS LIMITED TO THE REPAIR OR REPLACEMENT OF THE DEFECTIVE PARTS. GF SHALL NOT BE LIABLE FOR AND HEREBY DISCLAIMS ANY DIRECT, SPECIAL, INDIRECT, INCIDENTAL, EXEMPLARY OR CONSEQUENTIAL DAMAGES, INCLUDING, BUT NOT LIMITED TO: DAMAGES FOR LOSS OF PROFITS OR INCOME, LOSS OF USE, DOWNTIME, COVER, OR EMPLOYEE OR INDEPENDENT CONTRACTOR WAGES, PAYMENTS AND BENEFITS. CERTAIN STATES MAY CONFER ADDITIONAL RIGHTS REGARDING WARRANTIES AND IN THOSE STATES GF'S LIABILITY AND THE LIABILITY OF GF'S SUPPLIERS, SHALL BE LIMITED TO THE FULLEST EXTENT PERMITTED BY LAW.

The warranties contained herein, together with GF's current Terms and Conditions, contain all the representations and warranties with respect to the subject matter of this document, and supersede all prior negotiations, agreements and understandings with respect thereto. The recipient of this document hereby acknowledges and represents that it has not relied on any representation, assertion, guarantee, warranty, collateral contract or other assurance, except those set out in this document.

For additional information on this product or this warranty, please contact a GF Customer Service Representative.

NOTES:

1. Additional terms and conditions may apply. See GF's General Terms and Conditions on its website: www.grahamfield.com.
2. Freight claims must be notated on the appropriate shipping documents and must be made with immediacy. International, federal and state regulations govern specific requirements for freight claims. Failure to abide by those regulations may result in a denial of the freight claim. GF will assist you in filing the freight claim.
3. Claims for any short shipment must be made within three (3) days of the invoice date.

10 COMPLIANCE

The NSD system is a 24VDC powered add-on system for the healthcare facility transport stretchers.

The essential requirements of EC Machinery Directive 2006/42/EC have been applied and fulfilled.

The following applicable safety standards have been applied and fulfilled:

- EN-IEC 60204-1 Safety of machinery - Electrical equipment of machines - Part 1: General requirements
- EN-ISO 13849-1 Safety of machinery - Safety-related parts of control systems - Part 1: General principles for design

The following applicable requirements of EMC Directive 2004/108/EC have been applied and fulfilled:

- EN61000-6-1 Electromagnetic compatibility (EMC) - Part 6-1: Generic standards - Immunity for residential, commercial and light-industrial environment
- EN61000-6-3 Electromagnetic compatibility (EMC) - Part 6-3: Generic Standards - Emission standard for residential, commercial and light-industrial environment.

The relevant technical documentation has been compiled in accordance with Annex VII, Part B of EC Machinery Directive 2006/42/EC.

The partly completed machinery must not be put into service until the final machinery into which it is to be incorporated has been declared in conformity with the provisions of the Machinery Directive.

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Hausted® by graham-field

gf™ Proudly Made in the USA

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