840

user manual





SETUP
VIDEO

OPERATION
VIDEO

USER
MANUAL

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APPLICABLE MODEL(S)

This manual is for the following product numbers and serial numbers:

PRODUCT NUMBER	DESCRIPTION	EFFECTIVE S/N
840-003-22	840 Battery Powered Stretcher Bed, Mobile Base, 100-240V REAR Steer (rear swivel casters, front LH steering caster)	200101-XXXXXX to Current
840-003-23	840 Battery Powered Stretcher Bed, Mobile Base, 100-240V FRONT Steer (front swivel casters, rear RH steering caster)	200101-XXXXXX to Current
840-003-24	840 Battery Powered Stretcher Bed, Mobile Base, 100-240V SELECTIVE Steer (front LH & rear RH steering casters)	200101-XXXXXX to Current

REVISION HISTORY TABLE

DATE	REVISION	CO#/ECO#	DETAILS
9/8/2020	A	CO-0309/EC-0577	Initial release
12/18/2020	B	CO-0322	Added Effective Serial Numbers and created 840 In-service form
10/5/2021	C	CO-0361/EC-0634	Modified Hand Control Cord Section. Minor Text edits. Updated logo. Added QR code.
7/26/2022	D	CO-0385	Revised Set-up Instructions - modified step 2, added step 13.
12/10/2025	E	CO-0594	Updated images with new headrest knob. Added clarifying information in the Controls section about Momentary and Maintain Mode. Modified Accessories Table. Modified formant and In-Service Checklist. Changed Max Stack Height image to 2.
08/20/2026	F	CO-0675	Update to current branding standards. See CO-0675 for details.

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User Manual

MTI 840

GENERAL INFORMATION

Purpose of this Manual

This manual covers the safety, installation, instructions for use, and maintenance of this equipment. It is intended that it be used by the service professional installing the equipment and by the medical professionals who have been trained to use it. No repair information is included with this manual because no operator repairs are intended.

Intended Use

This equipment is intended to be used as a stretcher bed for general examinations and procedures performed by a qualified medical professional in a medical office environment.

Training Options

Training is based on the intended use of the equipment and the proficiency of the medical professionals who will be using the equipment. Training is accomplished either through reading the installation and operation sections of the user manual or through remote training provided by MTI Technical Support. To request training, contact MTI.

PRODUCT INFORMATION

Date of Purchase:_____

Model Number:_____

Serial Number:_____

(see "EQUIPMENT FEATURES" on page 10 for location of serial number label)

Symbols and Warnings

This manual contains Notices, Cautions, and Warnings, which are meant to call attention to particular parts or sections of this manual. Since safety is of our utmost concern, this manual should be read by all personnel who will be operating this medical equipment.

Symboles et Avertissements

Ce manuel contient des avis, des précautions et des avertissements qui sont destinés à attirer l'attention sur des pièces ou des sections de ce manuel particulier. Parce que la sécurité est notre première préoccupation, ce manuel doit être lu par tout le personnel qui sera d'utiliser cet équipement médical.

⚠ WARNINGS/AVERTISSEMENTS

Are used to call attention to condition, practice, or procedure, which could cause personal injury if not followed correctly. Please become familiar with all warnings and observe them at all times.

Sont utilisés pour attirer l'attention sur une condition, pratique ou procédure qui pourrait causer des blessures s'il n'est pas suivi correctement. S'il vous plaît se familiariser avec tous les avertissements et les observer à tout moment.

⚠ CAUTIONS/PRÉCAUTIONS

Are used to call attention to a condition, practice, or procedure which could cause product damage if not used correctly.

Sont utilisés pour attirer l'attention sur une condition, pratique ou procédure qui pourrait causer des dommages au produit s'il n'est pas utilisé correctement.

❗ NOTICES/AVIS

Are used to call attention to helpful information regarding a condition, practice, or procedure.

Sont utilisés pour attirer l'attention sur des informations utiles concernant une condition, pratique ou de procédure.

	*Type B applied part* *Type B, partie appliquée*
	*Proper shipping orientation for the product* *Orientation d'embarquement correct pour le produit*
	*Product is fragile; be careful in handling* *Le produit est fragile. Soyez prudent lors de la manipulation*
	*Protective earth ground* *Terre électrique de protection*
	*Alternating current* *Le courant alternatif*
	*Product must be kept dry* *Le produit doit rester sèche*
	*Upper and lower limits of temperature* *Limites haute et basse de la température*
	*Maximum stacking height of palletized units.* *Hauteur maximale de la pile d'unités sur palettes.*
	*Indicates the presence of a dangerous voltage/shock hazard* *Indique le présence de voltage dangereux/ de choc*
	*Follow instructions for use* *Suivez les instructions pour l'utilisation*
	*Consult operating instructions* *Consultez les instructions d'utilisation*
	*Manufacturer of product* *Fabricant de produits*
	*Date on which the product was manufactured* *Date de fabrication*
	*Identifies the manufacturer's serial number* *Identifie le numéro de série du fabricant*

	*Upper and lower limits of relative humidity* *Limites haute et basse de humidité relative*
	*Upper and lower limits of atmospheric pressure* *Limites haute et basse de la pression atmosphérique*
	*Location of fuses* *Emplacement de fusibles*
N	*Connection point for the neutral conductor* *Point de connexion pour le conducteur neutre*
L	*Connection point for the live conductor* *Point de connexion pour le conducteur sous tension*
G	*Connection point for the protective earth (ground) conductor* *Point de connexion pour le conducteur de protection (fil de masse)*
	*(WEEE) Waste of Electrical and Electronic Equipment* *Déchets d'équipements électriques et électroniques*
ETL CLASSIFIED c ETL us Intertek	*ETL listing mark* *Marque ETL Listé*
c UL us LISTED	*UL listing mark* *Marque UL Listé*
CE	*European Conformity mark* *Marque Conformité Européenne*
RoHS	*RoHS Compliant* *Conforme à RoHS*
	*Hospital Grade (green dot)* *De qualité hôpital (point vert)*
	*TUV listing mark* *Marque TUV Listé*
	*Emergency Stop* *Arrêt d'urgence*

	*Steering Mode (green)* *Mode de Direction (vert)*
	*Free Mode (yellow)* *Mode Libre (jaune)*
	*Locked Mode (red)* *Mode verrouillé (rouge)*

Safety Instructions

⚠ WARNING/AVERTISSEMENT

Read this manual before installation and use.

Lisez ce manuel avant l'installation et l'utilisation.

⚠ WARNING/AVERTISSEMENT

No modification of this equipment is allowed.

Aucune modification de cet équipement est autorisé.

⚠ WARNING/AVERTISSEMENT

Do not operate equipment in the presence of flammable anesthetics.

Ne pas faire fonctionner l'équipement en présence d'anesthésiques inflammables.

⚠ WARNING/AVERTISSEMENT

Do not leave the equipment unattended during use of programmable buttons (Home/P1-P4) or manual actuation of a function.

Ne laissez pas l'appareil sans surveillance pendant l'utilisation des positions programmées (position retournée/P1-P4) ou l'actionnement manuel d'une fonction.

⚠ WARNING/AVERTISSEMENT

Ensure equipment, patient, and other objects are in a safe condition before operating the equipment.

Assurer l'appareil, patient, et d'autre objets sont dans un état sûr avant d'utiliser l'appareil.

⚠ WARNING/AVERTISSEMENT

Verify patient arms, headrest and other adjustable equipment features and accessories are locked and in a safe and secure position before use.

Vérifiez que les accoudoirs, l'appui-tête et les autres éléments et accessoires de l'équipement ajustable sont verrouillés et dans une position sûre avant toute utilisation.

⚠ WARNING/AVERTISSEMENT

Ensure controls, cords, and other parts of the equipment do not create a tripping or other safety hazard for operator or patient.

Vérifiez les contrôles, les cordons et les autres parties de l'équipement ne créent pas un risque de trébuchement ou d'autres risques d'accident pour l'opérateur ou le patient.

⚠ WARNING/AVERTISSEMENT

To prevent unintentional movement of the equipment caused by accidental activation of the equipment controls, electromagnetic interference, or radio frequency interference from other equipment, use the On/Off Button (only on select models) on the hand control or hand switches. For equipment without the On/Off Button, use the alternate security method by activating the Safety Lockout setting. See "Safety Lockout" on page 15. Failure to do so may result in personal injury. Using the On/Off Button or the Safety Lockout setting will not prevent non-powered motion.

Pour éviter tout mouvement involontaire de l'équipement dû à une activation accidentelle des commandes, à des interférences électromagnétiques ou à des interférences radiofréquences provenant d'autres appareils, utilisez le bouton Marche/Arrêt (disponible sur certains modèles) de la commande manuelle ou des interrupteurs manuels. Pour les équipements dépourvus de bouton Marche/Arrêt, utilisez la méthode de sécurité alternative en activant le verrouillage de sécurité «Safety Lockout». Voir «Safety Lockout» à la page 15. Ne pas le faire peut entraîner des blessures corporelles. L'utilisation du bouton Marche/Arrêt ou du dispositif de verrouillage de sécurité n'empêchera pas les mouvements non motorisés.

⚠ WARNING/AVERTISSEMENT

Exceeding the specified load limit of any equipment component may result in personal injury or property damage. See "Specifications" on page 33.

Le dépassement de la limite de poids spécifiée de tout composant de l'équipement peut entraîner des blessures ou des dommages matériels. Voir «Specifications» à la page 33.

⚠ WARNING/AVERTISSEMENT

Backrest, seat, footrest, armrest, headrest, and other parts of the equipment should only be used for their intended purpose. MTI is not responsible for use of components outside of their intended use.

Le repose-dos, reste de siège, repose pieds, accoudoirs, appuie-tête et d'autres parties de l'équipement ne doivent être utilisés conformément à leur utilisation prévue. MTI n'est pas responsable de l'utilisation des composants au dehors de leur utilisation prévue.

⚠ WARNING/AVERTISSEMENT

During transport, lower the lift of the stretcher bed sufficiently to

prevent instability during transport.

Pendant le transport, abaissez suffisamment l'élévateur de la civière pour éviter toute instabilité.

⚠ WARNING/AVERTISSEMENT

This product can expose you to chemicals including nickel, which is known to the State of California to cause cancer. For more information go to www.P65Warnings.ca.gov and see www.mti.net/faqs for details on California Proposition 65.

Ce produit peut vous exposer à des agents chimiques, y compris le nickel, identifiés par l'État de Californie comme pouvant causer le cancer. Pour de plus d'informations, prière de consulter www.P65Warnings.ca.gov et voir www.mti.net/faqs pour les détails sur le Proposition de Californie 65.

Electromagnetic Interference

This equipment is designed and built to minimize electromagnetic interference with other devices. However, if interference is noticed between another device and this equipment, do one or more of the following:

1. Ensure other device is operating within its specifications.
2. Plug equipment into isolated circuit.
3. Remove interfering device from room.
4. Increase separation between equipment and interfering device.
5. Contact MTI if electromagnetic interference persists.

Electromagnetic Compatibility (EMC)

⚠ WARNING/AVERTISSEMENT

Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.

L'utilisation d'accessoires, de transducteurs et de câbles autres que ceux spécifiés ou fournis par le fabricant de cet équipement peut induire une augmentation des émissions électromagnétiques ou une diminution de l'immunité électromagnétique de cet équipement et entraîner un mauvais fonctionnement.

⚠ WARNING/AVERTISSEMENT

During use, portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the stretcher bed, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

Les équipements de communication RF portables (y compris les périphériques tels que les câbles d'antenne et les antennes externes) ne doivent pas être utilisés à moins de 30 cm (12 po) de toute partie de la civière, y compris les câbles spécifiés par le fabricant, pour éviter tout risque de dégradation des performances de cet équipement.

⚠ WARNING/AVERTISSEMENT

Use of this equipment adjacent to other electrical equipment should be avoided since it could result in improper operation. If such use is necessary, verify normal stretcher bed operation before use with patients.

L'utilisation de cet équipement à proximité d'autres équipements électriques doit être évitée car cela pourrait entraîner un mauvais fonctionnement. Si une telle utilisation est nécessaire, vérifier le fonctionnement normal de la civière avant de l'utiliser avec les patients.

❗ NOTICE/AVIS

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 normally 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 re-orienting the equipment.

Les caractéristiques d'émission de cet équipement le rendent approprié pour une utilisation dans les zones industrielles et les hôpitaux (CISPR 11 classe A). S'il est utilisé dans un environnement résidentiel (pour lequel la norme CISPR 11 de classe B est normalement requise), cet équipement peut ne pas offrir une protection adéquate aux services de communication radiofréquence. L'utilisateur peut avoir besoin de prendre des mesures d'atténuation, telles que le déplacement ou la réorientation de l'équipement.

This stretcher bed has been tested to comply with the requirements of EN 60601-1-2:2014 Ed.4 for electromagnetic compatibility (EMC). The stretcher bed was designed to be operated within the electromagnetic environment specified below.

Guidance and manufacturer's declaration - electromagnetic emissions		
Emissions Test	Compliance	Electromagnetic Environment - Guidance
RF emissions CISPR 11	Group 1	The 840 Stretcher Bed uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class A	The 840 Stretcher Bed is suitable for use in all establishments other than domestic and those directly connected to public low-voltage power supply network that supplies buildings used for domestic purposes.
Harmonic Emissions IEC 6100-3-2	Class A	
Voltage fluctuations/flicker emissions IEC 61000-3-3	Complies	

RECOMMENDED SEPARATION DISTANCES BETWEEN PORTABLE AND MOBILE RF COMMUNICATIONS EQUIPMENT AND THE MTI STRETCHER BEDS

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

Rated maximum output power of transmitter Watts (W)	Separation distances according to frequency of transmitter meters (m)		
	150 kHz to 80 MHz d = (1.17)(√P)	80 MHz to 800 MHz d = (1.17)(√P)	800 MHz to 2.5 GHz d = (2.33)(√P)
0.01	0.12	0.12	0.23
.1	0.37	0.37	0.74
1	1.17	1.17	2.33
10	3.69	3.69	7.38
100	11.70	11.70	23.30

For transmitters rated at a maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer.

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

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

GUIDANCE AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC IMMUNITY			
The MTI Stretcher Bed is intended for use in the electromagnetic environment specified below. The customer or the user of the MTI Stretcher Bed should assure that it is used in such an environment.			
Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment Guidance
Conducted RF IEC 61000-4-6 Radiated RF IEC 61000-4-3	3V rms 150kHz to 80 MHz 3V/m 80MHz to 2.7 Ghz	3V 3V/m	Portable and mobile RF communications equipment should be used no closer to any part of the MTI Stretcher Bed, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: d = (1.17)(√P) 150 kHz to 80 MHz d = (1.17)(√P) 80 MHz to 800 MHz d = (2.33)(√P) 800 MHz to 2.5 GHz Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and d is the recommended separation in meters (m). Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey [a] should be less than the compliance level in each frequency range. [b] Interference may occur in the vicinity of equipment marked with the following symbol: 
NOTE 1: At 80 MHz and 800 MHz, the higher frequency range applies. NOTE 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.			

NOTICE/AVIS

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

[a]Les forces de champ des émetteurs fixes, telles que les stations de base pour les téléphones radio (cellulaires / sans fil) et les radios mobiles terrestres, la radio amateur, les émissions de radio AM et FM et les émissions de télévision ne peuvent pas être prédites avec précision. Pour évaluer le environnement électromagnétique dû aux émetteurs RF fixes, une étude de site électromagnétique doit être envisagée. Si le mesuré la résistance au champ dans le lieu où le fauteuil de procédure MTI est utilisé dépasse le niveau de conformité RF applicable indiqué ci-dessus, le MTI Le président de procédure doit être observé pour vérifier le bon fonctionnement. Si des performances anormales sont observées, des mesures supplémentaires peuvent être nécessaires, telles que la réorientation ou le déplacement du lit de civière MTI.

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

GUIDANCE AND MANUFACTURER'S DECLARATION - ELECTROMAGNETIC IMMUNITY			
The MTI Stretcher Bed is intended for use in the electromagnetic environment specified below. The customer or the user of the MTI Stretcher Bed should assure that it is used in such an environment.			
Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment Guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 8 kV air	± 8 kV contact ± 2, 4, 8, 15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast Transient / burst IEC 61000-4-4	± 0.5, 1, 2 kV for power supply lines ± 1 kV for input/ output lines	± 0.5, 1, 2 kV for power supply lines ± 1 kV for input/ output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	± 0.5, 1 kV line(s) to line(s) ± 0.5, 1, 2 kV lines(s) to earth	± 0.5, 1 kV ± 0.5, 1, 2 kV	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11 Tested at 230, 220, 127 and 120 VAC.	0% U_T (100% dip in U_T) for 0.5 cycle 0% U_T (100% dip in U_T) for 1 cycle 70% U_T (30% dip in U_T) for 25 cycles 0% U_T (100% dip in U_T) for 250 cycles	0% U_T (100% dip in U_T) for 0.5 cycle 0% U_T (100% dip in U_T) for 1 cycle 70 % U_T (30 % dip in U_T) for 25 cycles 0% U_T (100% dip in U_T) for 250 cycles	Mains power quality should be that of a typical commercial or hospital environment. If the user of the MTI Stretcher Bed requires continued operation during power mains interruptions, it is recommended that the MTI Stretcher Bed be powered from an uninterruptible power supply or a battery.
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m, 50 / 60 Hz	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
NOTE: U_T is the a.c. mains voltage prior to application of the test level.			

Electrical Requirements

⚠ WARNING/AVERTISSEMENT

To avoid risk of electrical shock, this equipment must be connected to a supply mains with protective earth when the equipment is plugged in to a mains AC outlet.

Pour éviter tout risque de choc électrique, cet appareil doit être branché sur une prise électrique avec la terre électrique de protection, lorsque cet appareil est branché sur une prise électrique C.A.

⚠ WARNING/AVERTISSEMENT

When this equipment is plugged into a mains AC outlet, grounding reliability can only be achieved when the equipment is connected to an equivalent receptacle marked "Hospital Only" or "Hospital Grade".

Lorsque cet appareil est branché sur une prise électrique C.A., mise à la terre fiabilité ne peut être atteint lorsque l'appareil est connecté à un réceptacle équivalent marqué "Hôpital Uniquement" ou "de Qualité Hôpital".

⚠ WARNING/AVERTISSEMENT

When using endocardial catheters or high frequency surgical/ procedure devices, use non-conductive material to insulate the patient from metal portions of the equipment. Failure to comply may result in electric shock or burns to patient.

Lors de l'utilisation de cathéters endocavitaires ou dispositifs chirurgicale / procédure haute fréquence, utiliser un matériau non conducteur pour isoler le patient des parties métalliques de l'équipement. Le non-respect peut entraîner un choc électrique ou de brûlures pour le patient.

⚠ WARNING/AVERTISSEMENT

Equipment must be positioned in normal use such that the power cords can be easily accessed to unplug.

L'appareils doit être positionné en utilisation normale telle que les cordons d'alimentation peuvent être facilement accessibles à débrancher.

⚠ WARNING/AVERTISSEMENT

Equipment must be connected to an appropriate power source when loss of power source would result in an unacceptable risk.

L'appareils doivent être relié à une source d'alimentation appropriée en cas de perte de la source d'alimentation entraînerait un risque inacceptable.

⚠ WARNING/AVERTISSEMENT

Do not use a three to two prong adapter to plug the equipment into a two conductor outlet, as grounding reliability is not maintained.

Ne pas utiliser un adaptateur non de prise de terre pour brancher l'appareil sur une prise à deux conducteurs, la fiabilité de la terre n'est pas maintenue.

⚠ WARNING/AVERTISSEMENT

Do not plug this equipment into a low power or auxiliary outlet. This equipment should be connected to a 100-240 Volt AC (VAC), 47-63 Hertz (Hz) outlet. If this equipment is connected to an insufficient power source, it will cause irreparable damage to this device's electronics.

Ne branchez pas cet équipement sur une prise de faible puissance ou une prise auxiliaire. Cet équipement doit être branché sur une prise de 100-240 Volt AC (VAC), 47-63 Hertz (Hz). Si cet équipement est connecté à une source d'alimentation insuffisante, il causera des dommages irréparables à l'électronique de cet appareil.

! NOTICE/AVIS

When the equipment is powered solely by AC power, the power cord may be used to simultaneously isolate both poles of the power supply by removing the power cord at the equipment input or wall receptacle.

Lorsque l'équipement est alimenté uniquement par l'alimentation C.A., le cordon d'alimentation peut être utilisé pour isoler simultanément les deux pôles de l'alimentation en retirant le cordon d'alimentation à l'entrée de l'équipement ou la prise murale.

! NOTICE/AVIS

This equipment is compatible with electrosurgical generator devices. These devices may be used without degradation of the stretcher bed performance.

Cet équipement est compatible avec les générateurs électrochirurgicaux. Ces dispositifs peuvent être utilisés sans dégradation des performances de la civière mobile.

Contraindications

None.

Transportation and Storage Conditions

	Ambient Temperature	-10°C to +50°C (+14°F to +122°F)
	Relative Humidity	20-90% (non-condensing)
	Atmospheric Pressure	700 hPa to 1060 hPa (0.69 atm to 1.05 atm)

Disposal of Equipment

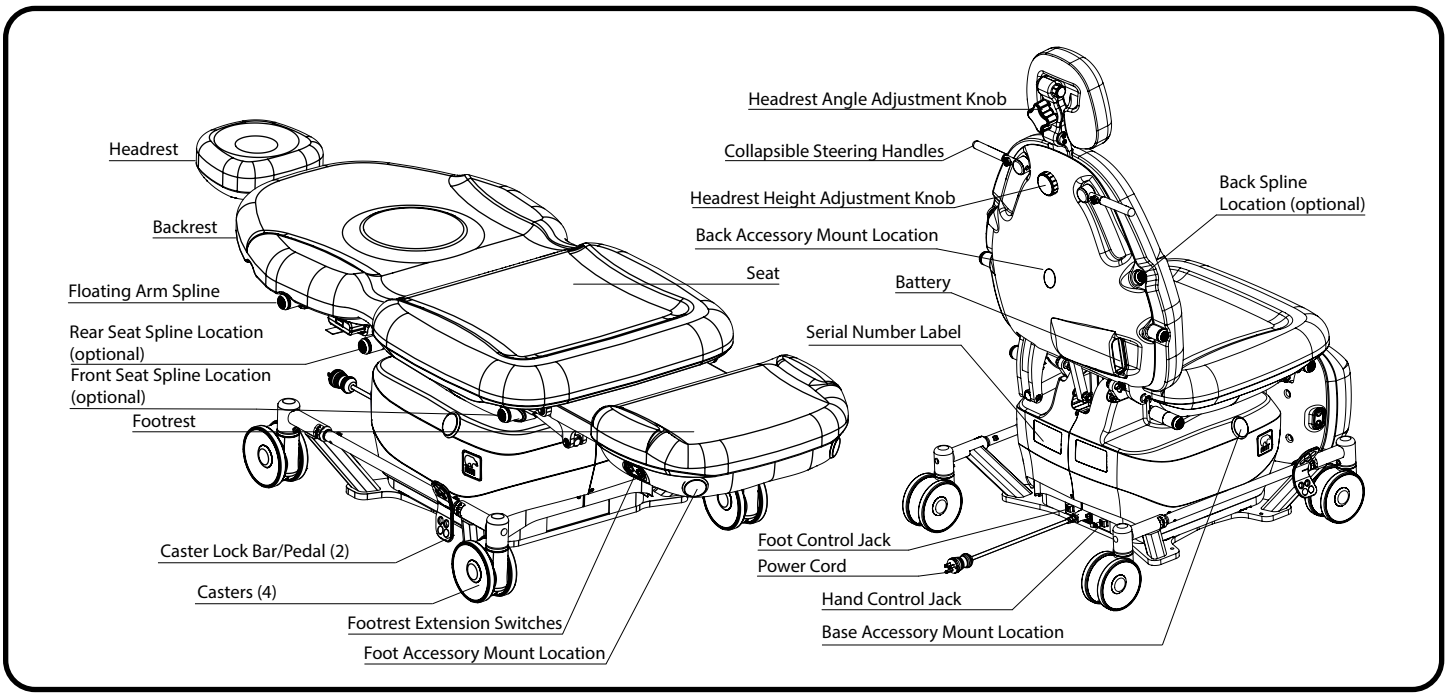
At the end of product life, the equipment, accessories, and other consumable goods may have become contaminated due to normal medical use. Consult local codes and ordinances for proper disposal of equipment, accessories and other consumable goods.

EQUIPMENT FEATURES

Description of 840 Mobile Battery Powered Stretcher Bed

The 840 Stretcher Bed has power lift, back, tilt, foot, and foot extension functions. It has a 725 lb. (328.8 kg) patient capacity. The 840 Stretcher Bed is 25” (63.5 cm) wide with an entry height of 18 3/4” (47.6 cm). It features a mobile base and collapsible steering handles for convenient maneuverability. It is powered by either mains AC power or a rechargeable, lithium ion battery. The bed will rise to a height of 37” (94 cm).

840 MOBILE BATTERY POWERED STRETCHER BED



INSTALLATION

⚠ WARNING/AVERTISSEMENT

This equipment is heavy. Get help to remove it from the shipping pallet. Use proper lifting procedures when lifting to avoid serious back injury.

Cet équipement est lourd. Obtenir de l'aide pour le retirer de la palette d'expédition. Utilisez les procédures propre de levage pour le soulever pour éviter grave blessure au dos.

⚠ WARNING/AVERTISSEMENT

After equipment is installed, verify programmable positions (Home/P1-P4) are safe and known positions.

Après l'équipement est installé, vérifier les positions programmables sont des positions sûres et connues.

⚠ CAUTION/PRÉCAUTION

To move equipment, lift only from the equipment's main structure.

Pour déplacer l'appareil, soulevez seulement de la structure principale de l'appareil.

NOTICE/AVIS

If this product has been exposed to low temperatures, allow adequate time for it to warm to room temperature before putting it into service.

Si ce produit a été exposé à de basses températures, laisser suffisamment de temps pour lui permettre de se réchauffer à température ambiante avant de le mettre en service.

NOTICE/AVIS

Inspect the shipping boxes for visible damage. Record on bill of lading, photograph, report any damage to the carrier immediately, and contact MTI Technical Support. Request an inspection by the carrier if any concealed damage is noted.

Inspectez les boîtes d'expédition pour les dommages visibles. Fiche sur le connaissement, prendre une photo, et signalez immédiatement tout dommage au transporteur et contactez le support technique de MTI. Demander une inspection par le transporteur en cas de dommage caché est noté.

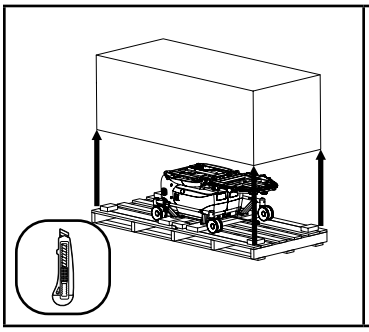
Required Tools

	Adjustable Wrench		Utility Knife		Flat Head Screw Driver		7/16" Wrench
	Set of Standard US Allen® Wrenches (for Accessories)		Phillips® Screwdriver (for Accessories)				

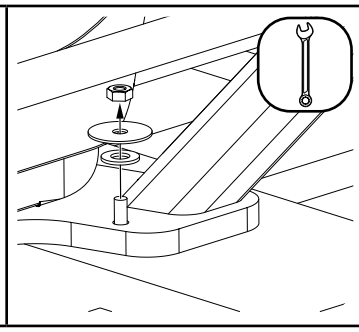
Explanation of Instructions

Instructions have steps which are designated by a number. Each step may also have sub-steps designated by a letter. For example, an instruction to fasten an accessory may have 4 steps (1-4) and each of those steps may have sub-steps (A-B) to be performed before moving onto the next step.

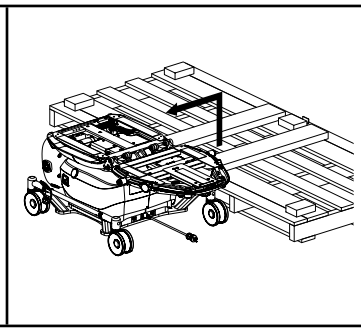
Unpacking and Set-up



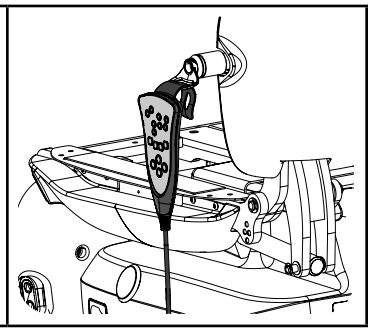
1. Remove shipping straps, shipping box, and other boxes from the pallet.



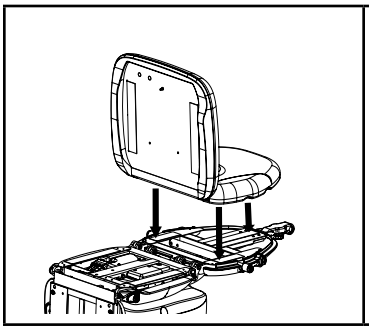
2. Remove the shipping bolts with an adjustable or 7/16" end wrench.



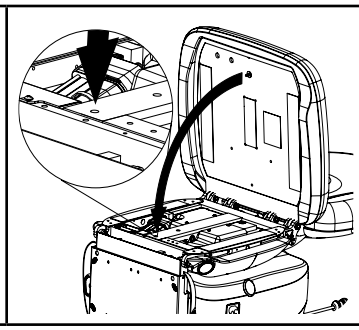
3. Slide the stretcher bed off the pallet. To move equipment, lift only from the equipment's main structure. Recycle packaging.



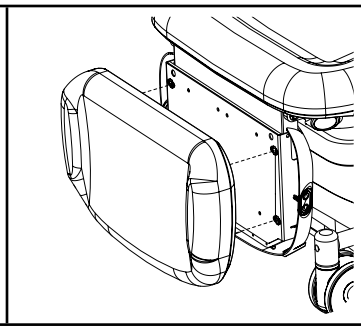
4. Temporarily install the Hand Control in the arm spline following the Hand Control instructions or set the hand control in safe place for later installation.



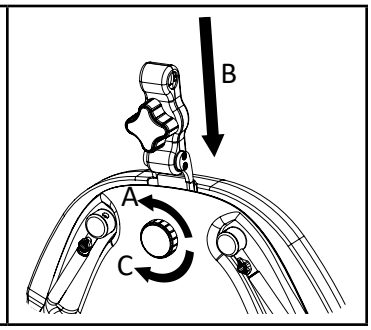
5. With the back and seat sections of the stretcher bed in the flat position, center the back portion of the seat/back cushion on the back section of the stretcher bed while holding up the seat portion of the cushion.



6. Lower the seat portion of the seat/back cushion into place and ensure that the ball stud enters the mating hole in the seat section of the stretcher bed.



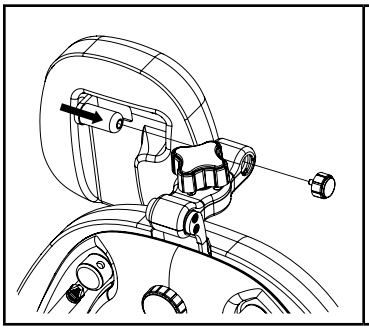
7. Attach the foot cushion by aligning the ball studs to the receptacles and pressing evenly to snap all 4 ball studs in place at the same time.



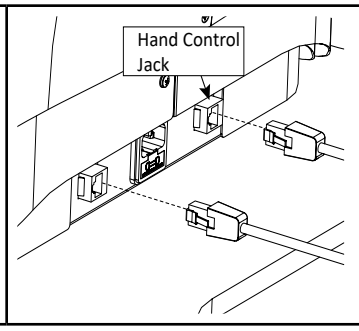
8. A. Loosen the knob on the back of the back section of the stretcher bed.

B. Insert the headrest lock mechanism into the slotted opening in the top of the backrest.

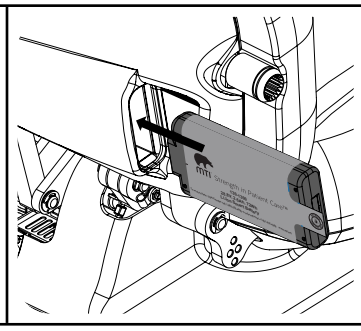
C. Tighten the knob on the backrest.



9. Loosen the knob on the articulating headrest mechanism and rotate it forward as shown. Remove the knob on the back of the headrest. Insert the headrest into lock mechanism. Reinsert knob, position headrest as desired, and tighten both knobs.

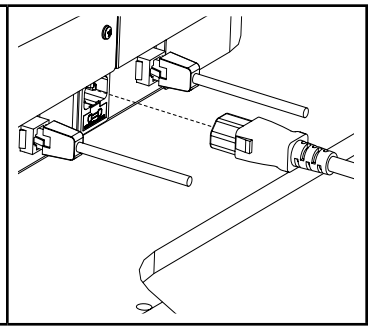


10. Plug in the hand control into the hand control jack on the patient's right hand side of the stretcher bed. If applicable, plug the foot control into the jack on the opposite side.



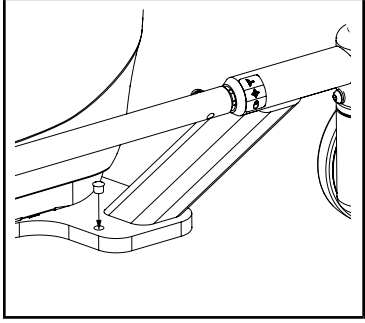
Battery Powered Operation

11. Note: Battery should be fully charged prior to use. See "Charging" on page 20. After charging, insert the battery into the holster in the back of the stretcher bed with the label facing out.

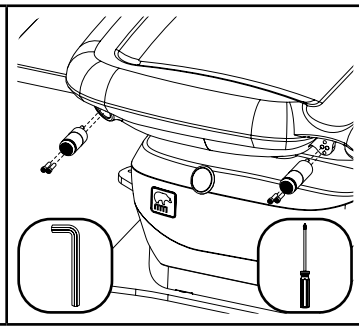


Mains AC Power Operation

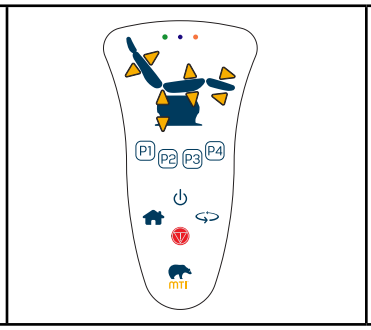
12. Insert the power cord at the base of the stretcher bed. Press firmly until it bottoms out. Plug the opposite end of the power cord into an appropriately marked receptacle.



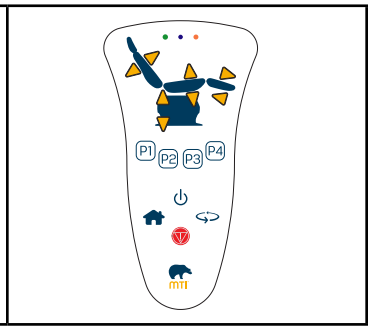
13. Insert the (4) provided hole plugs into the (4) holes that were used to attach the stretcher bed to the pallet.



14. If applicable, install any accessories (including the Hand Control if not previously installed). See the accessory's Instruction Guide for installation instructions.



15. Verify all functions of the stretcher bed are operational. Set the Smart Controller™ Settings for the Home and Program Buttons. See "Smart Controller™ Technology" on page 13 for instructions.



16. Save Home and Program Buttons' positions to desired locations. See "Controls" on page 15 for programming instructions.

OPERATION

Operating Conditions

	Ambient Temperature	+10°C to +40°C (+50°F to +104°F)
	Relative Humidity	30% to 75% (non-condensing)
	Atmospheric Pressure	700 hPa to 1060 hPa (0.69 atm to 1.05 atm)

Maximum surface temperature of applied part stretcher bed surfaces will not exceed 42°C.

SmartTech™

SmartTech™ is the combination of MTI's Smart Controller™ and Smart Safety™ technology using a centralized computer controller that enhances safety, reduces energy consumption, and provides smoother movement, additional memory options, and more user preferences.

Smart Controller™ Technology

The Smart Controller™ technology is the stack of technology features built into the stretcher bed, allowing the stretcher bed to move softly and smoothly with soft start/stop technology, a hibernation state to conserve power and a User Program Mode allowing users to change how some of the buttons function.

User Program Mode

The User Program Mode allows users to set how the Home, Program and Swivel Buttons function. For user preference these button settings can be changed by entering the User Program Mode to make the setting changes described below.

Home Button and Program Buttons Settings	
Momentary Button Setting	This setting is the factory default setting for the Home Button and Program Buttons. It allows the user to press and release the Home Button or Program Buttons to automatically move the stretcher bed to the button's saved position. Press any button, except the same button that was pressed to initiate movement, to stop automatic motion.
Maintain Button Setting	This setting allows the user to press and hold the Home Button or Program Buttons to move the stretcher bed to the button's saved position. Release the button to stop movement of the stretcher bed.

To enter the Smart Controller™ User Program Mode:

1. Press and Hold the **Stop** Button for 5 seconds. The stretcher bed will beep once and the blue and orange LEDs on the hand control will flash twice. The Foot Control does not have LEDs, but there are corresponding lights on the stretcher bed PCA inside the stretcher bed.
2. Press the following sequence of buttons (within 30 seconds from the time the Stop Button is released) to enter the Smart Controller™ User Program Mode: **Back Up, Back Up, Back Down, P2, Stop**. Each button press will beep and, if the correct sequence is entered, the stretcher bed will sound a low-high tune. If the wrong sequence of buttons is entered, the stretcher bed will sound three low beeps and return to normal operation.
3. Within a subsequent 30 seconds, press the following sequences of buttons to change the Home Button and Program Buttons Settings.

Home Button and Program Buttons Settings	
Momentary Button Setting	Back Up, Back Up, P1
Maintain Button Setting	Back Up, Back Up, P2

4. The stretcher bed will sound a high-low-high tune to accept the setting and the blue LED will flash twice. If the wrong buttons are pressed, the stretcher bed will sound three low beeps and the blue LED will not flash.
5. To exit the Smart Controller™ User Program Mode, press the Home Button. The stretcher bed will sound a high-low tune and only the green light will be on. The stretcher bed will also automatically exit the Smart Controller™ User Program Mode after 30 seconds if no buttons are pressed.
6. To confirm if the stretcher bed is in the desired setting (Momentary Button Setting or Maintain Button Setting), use the manual buttons (Lift Up or Tilt Up) to move the stretcher bed to a position away from the Home Position. Press and release the Home Button. If the stretcher bed moves automatically towards the Home Position, the stretcher bed is in the Momentary Button Setting. If the stretcher bed stops moving as soon as the Home Button is released, the stretcher bed is in the Maintain Button Setting.

Hibernation State

To conserve power, after 60 minutes of non-use the stretcher bed will automatically hibernate. To wake the stretcher bed from the Hibernation State, press any button.

Smart Safety™ Technology

The Smart Safety™ technology is the safety features built into the Smart Controller™, making the stretcher bed safer and easier to use. The Smart Safety™ technology includes the Collision Avoidance System™, which establishes Safe Zones that will disable a function(s) when the stretcher bed encounters a Safe Zone boundary. When the Collision Avoidance System™ detects a Safe Zone boundary, the stretcher bed will sound a single tone and disable the stretcher bed function(s). To re-enable full stretcher bed functionality, move the stretcher bed function into the Safe Zone.

Safety Lockout

The Safety Lockout disables all commands to prevent unwanted or inadvertent movements.

To enable the Safety Lockout, press the following sequence of buttons (within 1 second between presses): **Stop, Stop, Back Up, Stop.**

To disable the Safety Lockout, press the following sequence of buttons: **Stop, Stop, Back Down, Stop.**

Shutdown

To shutdown the 840 Stretcher Bed, press the On/Off Button on the hand control, or remove the battery from the stretcher bed and disconnect the power cord from the stretcher bed or the power outlet. See “Power Cord Removal Instructions” on page 27.

Controls

ITEM	DESCRIPTION	FUNCTION
FOOT AND HAND CONTROLS		
1	Lift Up/Down Buttons	Raises or lowers the stretcher bed top by pressing the desired direction.
2	Back Up/Down Buttons	Raises or lowers the backrest by pressing the desired direction.
3	Tilt Up/Down Buttons	Decreases or increases the angle of the seat by pressing the desired direction.
4	Foot Up/Down Buttons	Raises or lowers the footrest by pressing the desired direction. The footrest will automatically extend while pressing Foot Up and, when held, will continue to extend the foot fully. The Foot Down Button will automatically retract the foot as it approaches the floor.
5	Stop Button(Emergency Stop Button)	Stops all movement of stretcher bed.
6	Swivel Button	This button is non-functional on this stretcher bed.
7	Home Button To determine if the stretcher bed is in the Momentary Button Setting or the Maintain Button Setting see “User Program Mode” on page 14.	The Home Button moves the stretcher bed to either a Programmed Home Position or an ADA Compliant Home Position. The Programmed Home Position is user programmable and can be saved in any preferred position. **Momentary Button Setting (Default):** Press and release the Home Button to move the stretcher bed to the Programmed Home Position. Press and release the Home Button twice within 1 second to move the stretcher bed to the ADA Compliant Home Position (Lift Down, Tilt Down, Back Up, and Foot Down and Retracted). To save a new Programmed Home Position, position the stretcher bed to the desired position then press and hold the Home Button for 10 seconds. The stretcher bed will play a high-low-high tune after the Programmed Home Position has been saved. **Maintain Button Setting:** Press and hold the Home Button to move the stretcher bed to the Programmed Home Position. After arriving at Programmed Home, release the Home Button and then press it again within 1 second to move the stretcher bed to the ADA Compliant Home position (Lift Down, Tilt Down, Back Up, and Foot Down and Retracted). To save a new Programmed Home Position; position the stretcher bed to the desired position, press and release the Stop Button twice, then hold the Home Button for 10 seconds. The stretcher bed will play a high-low-high tune after the Programmed Home Position has been saved.

8	Program Buttons (P1, P2, P3, P4) To determine if the stretcher bed is in the Momentary Button Setting or the Maintain Button Setting see “User Program Mode” on page 14.	The Program Buttons move the stretcher bed to a user defined Program Position. NOTE: The stretcher bed has no pre-existing Program Positions upon arrival and will only beep and not move until Program Positions have been saved. The Program Buttons have two button settings that can be changed by the user. **Momentary Button Setting (Default):** Press and release to automatically move the stretcher bed to the Program Position. To save a Program Position, position the stretcher bed to the desired position then press and hold a Program Button for 5 seconds. The stretcher bed will play a high-low-high tune after the position has been saved. **Maintain Button Setting:** Press and hold the button to move the stretcher bed to the Program Position. To save a Program Position; position the stretcher bed in the desired position, press and release the Stop Button twice, then hold the Program Button for 5 seconds. The stretcher bed will play a high-low-high tune after the position has been saved.
9	On/Off Button	Turns the stretcher bed on and off (Conserves power and prevents accidental actuation of the stretcher bed).
10	LED Indicators	Used to provide user feedback on conditions, inputs, and errors. Left to right are green, blue, and orange LEDs. See “GUIDE TO INDICATOR LIGHTS AND SOUNDS” on page 31 for details.
FOOT EXTENSION SWITCHES		
11	Foot Extend/Foot Retract Buttons	Independently extends or retracts the foot section by pressing the desired direction. NOTE: Footrest will not extend until foot has been raised at least 20° and moved away from Safe Zone boundary. When foot is raised, footrest retraction is limited to prevent interference with seat.

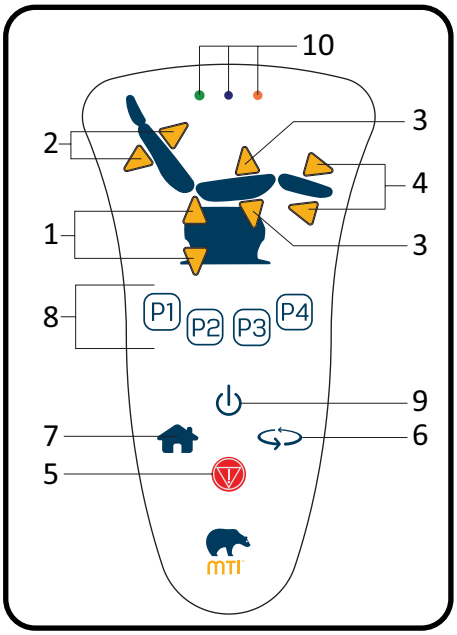
NOTICE/AVIS

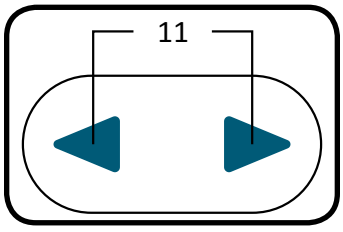
For additional safety, any button will stop all automatic motions except for the same button that was pressed to initiate movement.
Pour plus de sécurité, n’importe quel bouton peut arrêter tous les mouvements automatiques à l’exception du bouton même qui a été pressé de commencer le mouvement.

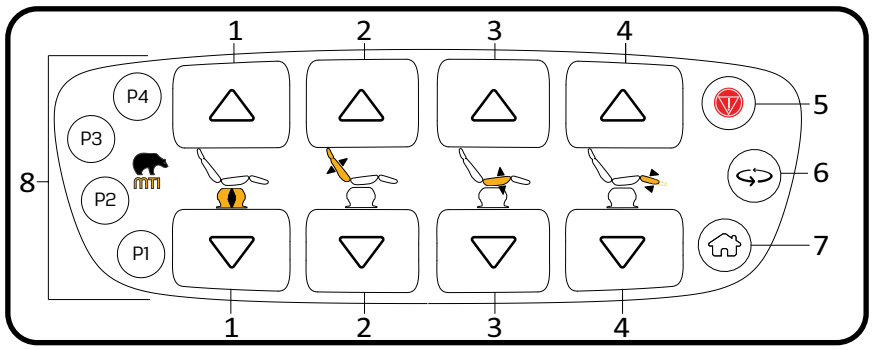
NOTICE/AVIS

Swivel Button is present on all foot and hand controls. For this stretcher bed, this button is not functional.

Le bouton de pivotement est présent sur tous les commutateurs au pied et à main. Pour cette civière, ce bouton n’est pas fonctionnel.

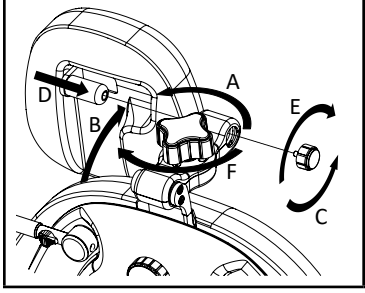






Headrest Operation

Articulating Headrests



1. To attach the headrest.

A. Loosen the knob on the articulating headrest lock mechanism.

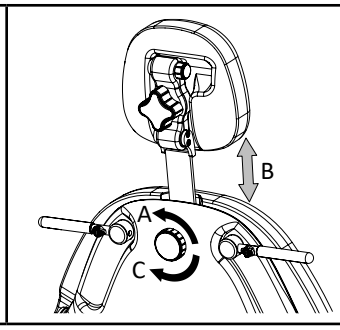
B. Rotate the articulating headrest lock mechanism forward.

C. Remove the small knob on the headrest pintle (turn counterclockwise).

D. Insert the headrest pintle into the articulating headrest lock mechanism.

E. Reattach the small knob into the end of the headrest pintle (turn clockwise).

F. Reposition the headrest to a desired location and tighten the knob on the articulating headrest mechanism.



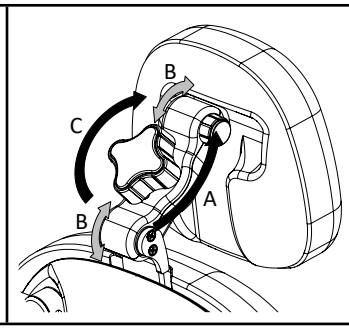
2. To adjust the height of the headrest:

A. To loosen, turn the knob on the back of the stretcher bed counterclockwise.

B. Adjust the height of the headrest.

C. To secure, turn the knob clockwise.

The headrest can be completely removed, if desired, when the knob is loosened.

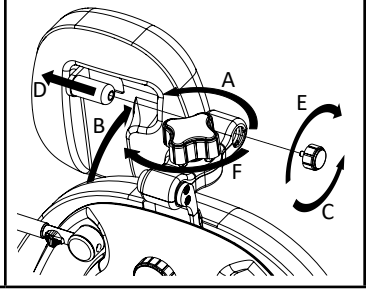


3. To adjust the angle of the headrest:

A. To loosen, turn the knob on the articulating headrest lock mechanism counterclockwise.

B. Adjust the position and angle of the headrest.

C. To secure, turn the knob clockwise.



4. To remove a headrest:

A. Loosen the knob on the articulating headrest lock mechanism.

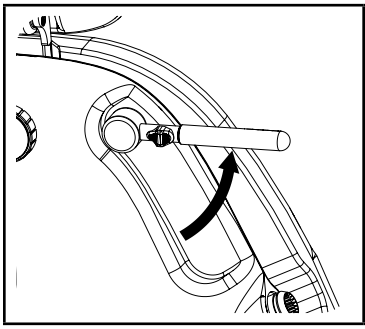
B. Rotate the articulating headrest lock mechanism forward.

C. Remove the small knob on the headrest pintle (turn counterclockwise).

D. Pull the headrest completely out of the headrest lock mechanism.

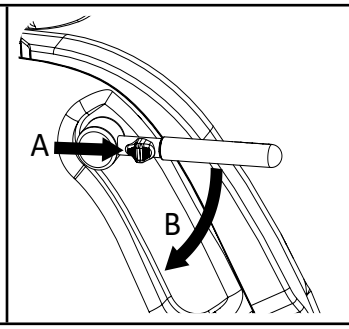
E. Reattach the small knob to the end of the headrest pintle to keep it from being lost (turn clockwise).

Steering Handle Operation



1. To raise the steering handle:

Rotate the handle upwards until the handle locks into a horizontal position.



1. To collapse the steering handle:

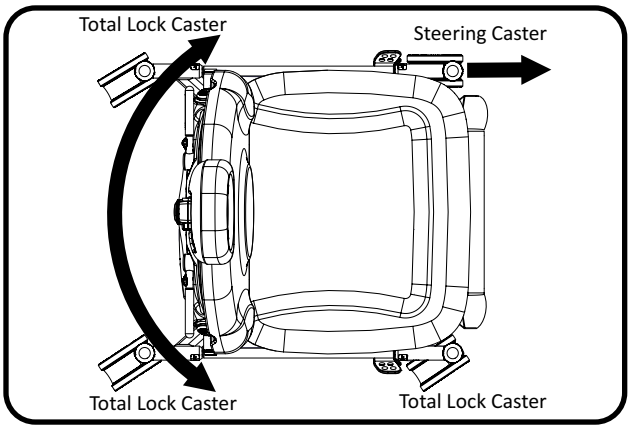
A. Slide and hold the handle release latch outward.

B. Lower the handle, release the latch and confirm the handle snaps into the lower position.

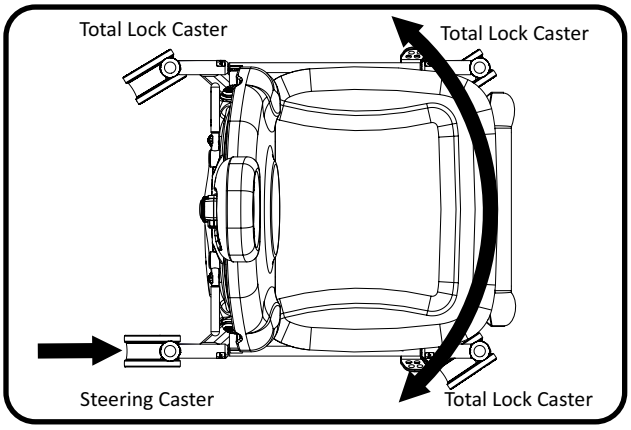
Mobile Base Operation

The 840 Stretcher Bed can be ordered in one of three different Steering Options. Each is indicated by a two digit number at the end of the stretcher bed's item number. E.g. XXX-XXX-22. See the Serial Number Label to determine

1. 840-003-22 is the REAR Steer Option (rear swivel casters, front LH steering caster) The Rear Steer Option utilizes the front caster on the patient's left side as the Steering Caster and the other casters are Total Lock Casters.
2. 840-003-23 is the FRONT Steer Option (front swivel casters, rear RH steering caster). The Front Steer Option utilizes the rear caster on the patient's right side as the Steering Caster and the other casters are Total Lock Casters.
3. 840-003-24 is the SELECTIVE Steer (front LH & rear RH steering casters) This option allows the user to choose if they want to Front Steer or Rear Steer. The Selective Steer Option utilizes both the front caster on the patient's left side and the rear caster on the patient's right side as the Steering Caster. The other casters are Total Lock Casters.



REAR STEER OPTION

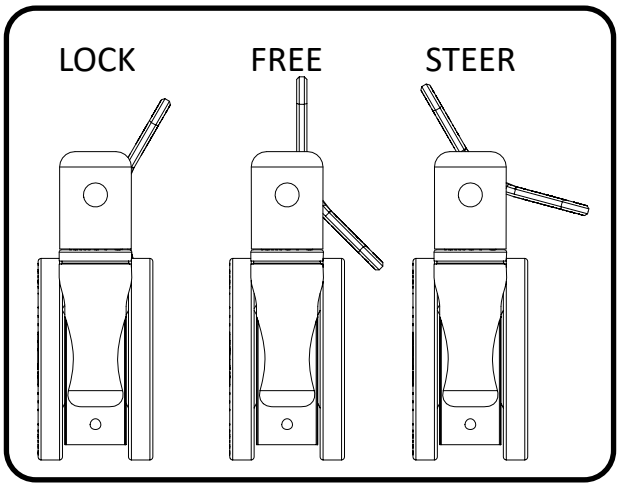


FRONT STEER OPTION

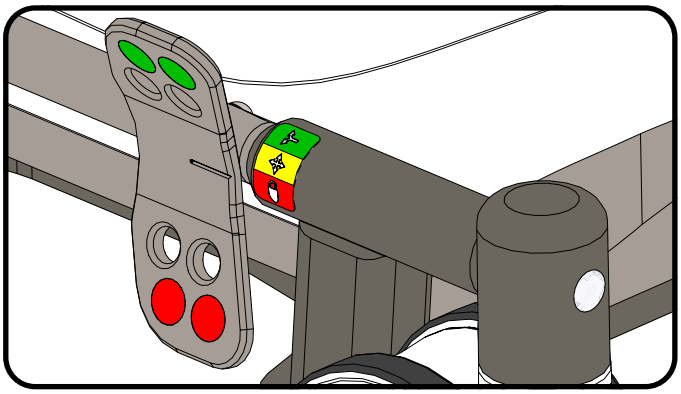
The locking bar has 3 positions. In the LOCK position, the Steering Caster will not roll or swivel. In the FREE position, the Steering Caster will swivel and roll. In the STEER position, the Steering Caster will roll but not swivel.

In the LOCK position, the Total Lock Casters will not roll or swivel. In the FREE or STEER position, the Total Lock Casters will roll and swivel.

To move the pedal from one mode to the other, firmly press down on the lower pedal or, with either the toe or heel of the shoe, lightly tap the top of the pedal until the pedal indicator line points to the desired mode on the indicator label.

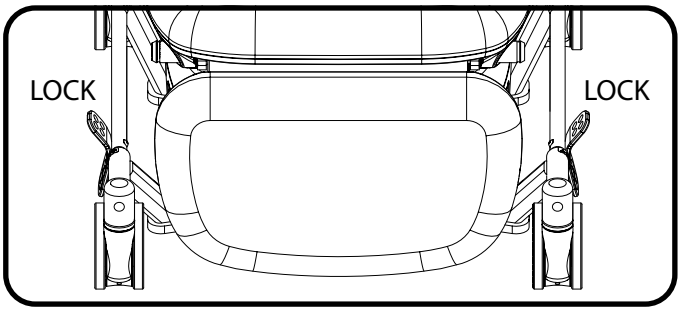


MODE INDICATOR: Each locking bar has a mode indicator line on the center of the pedal that points to the adjacent indicator label on the base of the stretcher bed. Note that the (3) Total Lock Casters operate the same whether they are in FREE or STEER Mode.



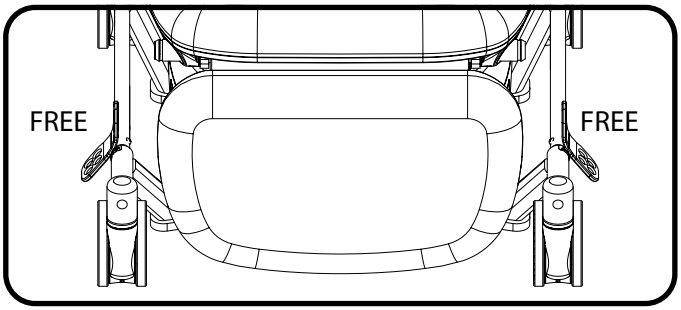
MODE INDICATOR

LOCK MODE: This mode is used to prevent movement of the stretcher bed. When both pedals are in the LOCK position; all (4) casters will not roll or swivel.



LOCK MODE

FREE MODE: This mode is used when you need to pull the stretcher bed away from an object in a perpendicular direction or turn the stretcher bed with a zero radius. When both pedals are in the FREE position; all (4) casters will roll and swivel freely.



FREE MODE

STEERING MODE: This mode is used for transport as the Steering Caster acts as a steering wheel to aid in easy transport. When one pedal is in the STEER position the opposite pedal should be in the FREE position; the Total Lock Casters will roll and swivel freely and the Steering Caster(s) will roll but will not swivel.

⚠ WARNING/AVERTISSEMENT

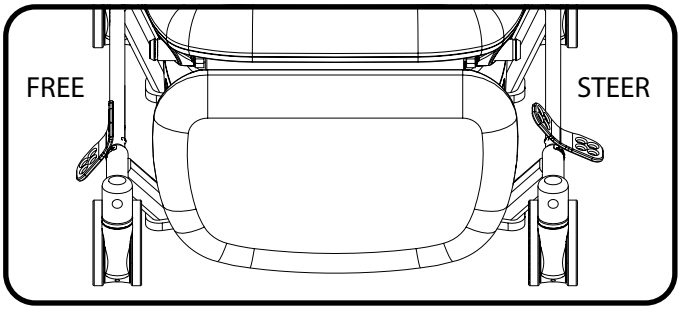
If the stretcher bed has the Selective Steer Option, do not put both the left-hand and right-hand locking bar in the STEER Position. This will prevent the stretcher bed from turning.

Si la civière mobile a l'option de Conduire Sélective, ne mettez pas la barre de verrouillage à droite et à gauche dans la position de STEER (CONDUIRE) à la fois. Cela interdire la civière mobile de tourner.

NOTICE/AVIS

Orient the steering caster opposite the direction of travel to prevent the casters from wobbling.

Orientez la roulette de direction opposée à la direction de voyage pour éviter le tremblement des roulettes.



STEER MODE

BATTERY & CHARGER OPERATION

NOTICE/AVIS

The battery does not charge while plugged into the stretcher bed even if the stretcher bed is plugged into an outlet. See charging instructions for directions on charging the battery.

La batterie ne se recharge pas lorsqu'il est branché dans le fauteuil même si la civière mobile est branchée sur une prise. Voir les instructions de chargement pour les directions sur la charge de la batterie.

Charging

The battery has an LCD fuel gauge located on the end of the battery. This fuel gauge displays five (5) bars when fully charged. Each bar represents twenty percent of the battery's capacity (Figure 1).

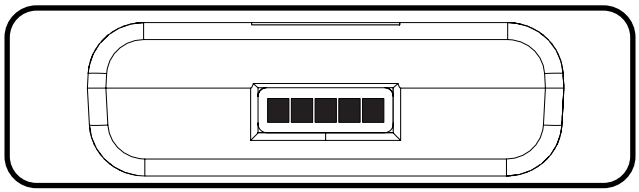


Figure 1

Fully charge battery prior to use. To charge the battery,

1. Plug the charger power cord into a properly grounded 100-240 VAC electrical outlet.
2. Place the charger on a flat surface away from heat and moisture.
3. Align the battery connector and charger connector and push them together. The MTI Logo on the battery and label pull tab should face out (Figure 2).
4. The charger will automatically begin to charge the battery. The LCD fuel gage on the battery will flash to indicate charging is in process.

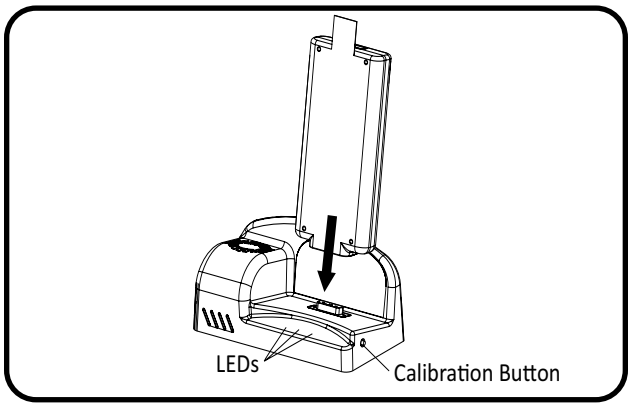


Figure 2

NOTICE/AVIS

MTI recommends the following procedure be followed in order to ensure that the battery is always ready for use and maintains a higher life expectancy.

MTI recommande de suivre la procédure suivante afin de s'assurer que la batterie est toujours prête à être utilisée et conserve une durée de vie plus longue.

MTI provides (2) batteries for each stretcher bed purchased. Best practices are as follows:

1. At the end of the day remove the battery from the stretcher bed and charge it.
2. Before the next surgery, insert a fully charged battery into the stretcher bed.
3. Alternate batteries every day.

NOTICE/AVIS

Leaving the battery on the charger, even when fully charged, will not damage the battery or the charger.

Le fait de laisser la batterie sur le chargeur, même lorsqu'il est complètement chargé, n'endommagera pas la batterie ou le chargeur.

Charge time for a fully depleted battery should typically be less than 2 hours, at which time the green LED on the charger should be solid and the fuel gauge on the end of the battery should show five bars.

LED Indication

The charging status of the battery is indicated by three LEDs on the front of the charger:

Green flashing	Battery is charging
Green solid	Battery is fully charged
Blue flashing	Battery is being calibrated
Blue Solid	Battery is calibrated
Red Flashing	Battery fuel gauge needs calibration
Red Solid	Error or battery is unrecoverable

Discharging

If the stretcher bed's power cord is plugged into a mains AC outlet, the bed will be powered by mains AC power, even with a battery inserted into the stretcher bed battery holster. If AC power is removed or unplugged, the stretcher bed will automatically switch to battery power.

⚠CAUTION/PRÉCAUTION

If the stretcher bed is not in use for 1 or more days and is not plugged into mains AC power, remove the battery from the stretcher bed, otherwise the battery will discharge. Leaving the battery in a non-AC powered stretcher bed for an extended period of time may result in premature failure of the battery.

Si la civière mobile n'est pas utilisée pendant 1 ou plusieurs jours et n'est pas branchée sur l'alimentation C.A., retirez la batterie de la civière mobile, sinon la batterie se déchargera. Laisser la batterie dans une civière mobile qui n'est pas branchée à l'alimentation C.A. pendant une période prolongée peut entraîner une panne prématurée de la batterie.

The runtime of the battery will be reduced if it is operated below room temperature.

Calibration

Frequent partial discharges will reduce the accuracy of the fuel gauge. The battery has a built-in monitoring system which checks the accuracy of the fuel gauge, based on the discharge history of the battery. A flashing red LED on the charger will indicate a battery fuel gauge needs calibration.

Calibration of the battery fuel gauge is achieved by plugging the charger into a properly grounded outlet, connecting the battery to the charger, and then pressing the calibration button on the side of the charger (See Figure 2). While calibrating the battery, keep the charger away from direct sunlight or heat sources in order to avoid calibration errors. There may be a short delay before the calibration begins.

Calibration may take 25-27 hours. The charger will fully charge, then fully discharge the battery to calibrate the fuel gauge, and then the battery will be fully recharged.

The battery and charger will get warm during the calibration procedure. The Blue LED will be solid when the calibration is complete.

Battery Life

End of life on the battery is defined as the point at which the battery fails to deliver 80% of its original rated capacity. This will occur after approximately 300 charge/discharge cycles.

Additional Warnings, Cautions and Notices

⚠ WARNING/AVERTISSEMENT

Do not expose the battery, charger, or power supply to liquids.

Ne pas exposer la batterie, le chargeur, ou l'alimentation à des liquides.

⚠ WARNING/AVERTISSEMENT

Do not expose to, or dispose of the battery in fire.

N'exposez pas et ne jetez pas la batterie au feu.

⚠ WARNING/AVERTISSEMENT

Do not short-circuit the battery.

Ne court-circuitez pas la batterie.

⚠ WARNING/AVERTISSEMENT

Only use MTI supplied batteries in the 840 Stretcher Bed.

Utilisez uniquement les batteries fournies par MTI dans la civière mobile 840.

⚠ WARNING/AVERTISSEMENT

Avoid excessive physical shock or vibration to the battery.

Évitez les chocs physiques excessifs ou les vibrations de la batterie.

⚠ WARNING/AVERTISSEMENT

Do not use a battery that is damaged.

N'utilisez pas une batterie endommagée.

⚠WARNING/AVERTISSEMENT

Do not open the battery, charger, or power supply case. There are no user serviceable parts inside.

N'ouvrez pas la batterie, le chargeur ou le boîtier d'alimentation. Il n'y a aucune pièce réparable par l'utilisateur à l'intérieur.

⚠CAUTION/PRÉCAUTION

Do not cover the charger case or leave in direct sunlight, as this could cause overheating.

Ne couvrez pas le boîtier du chargeur ou laisser à la lumière directe du soleil, car cela pourrait entraîner une surchauffe.

⚠CAUTION/PRÉCAUTION

If the battery is stored for an extended period of time, the battery must be fully charged every 6 months in order to avoid premature failure of the battery.

Si la pile est stockée pour longtemps, la batterie doit être chargée complètement tous les 6 mois pour éviter une défaillance prématurée de la pile.

⚠ CAUTION/PRÉCAUTION

If a battery is completely drained (no bars on fuel gauge) it must be immediately charged. If it is not charged, the battery will enter a shutdown mode. If the battery is left in shutdown mode, the battery will reach an unrecoverable state where it cannot be charged and must be replaced (not covered by warranty). A battery that is left in shutdown mode too long will be indicated by a solid red LED on the battery charger when it is connected.

Si une batterie est complètement déchargée (pas de barre sur la jauge de carburant), elle doit être chargée immédiatement. Si elle n'est pas chargée, la batterie passera en mode d'arrêt. Si la batterie est laissée en mode d'arrêt, elle atteindra un état irrécupérable où elle ne pourra pas être chargée et devra être remplacée (non couverte par la garantie). Une batterie laissée trop longtemps en mode arrêt est indiquée par un voyant rouge fixe sur le chargeur de batterie lorsqu'il est connecté.

⚠ CAUTION/PRÉCAUTION

The battery must be charged with MTI's charger. If another charger is used, battery damage or other unknown events may occur.

La batterie doit être chargée avec le chargeur de MTI. Si un autre chargeur est utilisé, les dommages de la batterie ou autres événements inconnus peuvent se produire.

❗ NOTICE/AVIS

Place the charger in a cool spot, away from external heat sources.

Placer le chargeur dans un endroit frais, à l'écart des sources de chaleur externes.

Storage

MTI recommends that the battery and charger be stored at room temperature. Elevated temperatures will reduce storage life. MTI lithium ion batteries are shipped partially to fully charged to give at least 6 months of shelf life at room temperature. However, it is recommended to fully charge the battery before the stretcher bed is operated or prior to storing the battery. It is recommended that the battery is recharged every 6 months if in storage.

Charger Specifications

Weight	11 oz / 308 grams
H x L x W	3.5 x 7 x 3.6"/88 x 180 x 92 mm
Energy Compliance	Compliant to California Energy Commission (CEC) Standard: DoE CFR, Title 10, Chapter II, Subchapter D, Part 430
Communications Compliances	System Management Bus Rev 1.0, Smart Battery Data Specification rev 1.0 and Smart Battery Charger specification Rev 1.0 compliant
Power Supply	100-240VAC, 36-38V, 1.73A DC, 150x35x70mm, 600g /21oz UL listed, CE Compliant
Mains Power (AC) Cord	110V N. American 3-pin connector

Battery Disposal

MTI lithium ion batteries are classified by the US federal government as non-hazardous waste and are safe for disposal in the normal municipal waste stream. However, the batteries do contain recyclable materials and are accepted for recycling by a number of regional battery recycling programs. MTI encourages you to recycle your batteries. The battery charger must be recycled and cannot be disposed of in a landfill.

In North America, contact the Rechargeable Battery Recycling Corporation (RBRC) at www.call2recycle.org.

❗ NOTICE/AVIS

The battery contains lithium ions. The amount of lithium ions contained in the battery are safe for consumer use.

La batterie contient des ions lithium. La quantité d'ions lithium contenus dans la batterie est sans danger pour les consommateurs.

LIST OF ACCESSORIES

DESCRIPTION	USE	RESTRICTIONS
Headrests		
Deep Oval, Horseshoe, Magnetic, Ophthalmic, Oval, Prosthetic, and Rectangular Articulating Headrest	Provides an articulating surface with various pillow shapes/sizes to support patient's head.	
Arms		
Arm, IV/Surgery, 18"Deluxe or Standard 8-way Seat Rail Mtd	Provides an articulating surface for patient's arms.	Requires Spline Assy, A Size (set of 2) mounted in Front Seat Spline Location. Requires Accessory Mtg Rail Assy.
Arm, IV/Surgery, 18" Deluxe or Standard 8-way Floating Mtd	Provides an articulating surface for patient's arms.	
Arm, IV/Surgery, 24" Deluxe 8-way Back Rail Mtd	Provides an articulating surface for patient's arms.	Requires Spline Assy, A Size (set of 2) mounted in Back Spline Location. Requires Accessory Mtg Rail Assy.
Arm, Patient, 4-way Floating	Used by patient to rest their arms during exams/procedures.	
Arms, Patient, Slide Back Floating (pair, LH, or RH)	Used by patient to rest their arms during exams/procedures.	
Procedure Table, Hand/Forearm	Provides an articulating surface for patient's arms.	
Lighting Options		
Adptr Kit, Back Mtd Exam Light w/100-240V Transformer	Provides a mount on the back of the stretcher bed and transformer for 12-24V Lumerus Exam Light.	Not compatible with Paper Roll Holder.
Exam Light, Lumerus LED, Flex Neck, 12-24V DC	Provides lighting during patient procedures.	Requires Adptr Kit, Back Mtd Exam Light w/100-240V Transformer.
Exam Light, Lumerus LED, Flex/Rigid Neck, 12-24V DC	Provides lighting during patient procedures.	Requires Mayo Tray w/Light Adptr or Post Assy w/Light Adptr.
Other Accessories		
Accessory Mount, Base	Provides a base mount for Mayo Trays, IV Rods, Lights, and Cardiac Monitor Trays.	
Accessory Mtg Rail Assy, 829 Seat & 830 Back/Seat	Provides standard hospital rails to attach to the seat or back of the stretcher bed for mounting accessories.	Requires Spline Assy, A Size (set of 2) mounted in Front Seat or Back Spline Locations.
Bed Rail & Transfer Support, ADA Compliant	Provides a support for entry and exit into the chair. Also used as flip-up bed rail to secure a patient.	Cannot be inserted while an arm is in the Floating Arm Spline. Requires Spline Assy, B Size (Set of 2) mounted in Rear Seat Spline Location.
Exam Boots, GStirrup w/Gel Pad Lithotomy (Pair)	Provides a pair of boots to stabilize the lower legs of a patient.	Requires Stirrups, Seat Spline Mtd (pair).
Flexible Oxygen Delivery System	Provide a flexible arm to deliver oxygen to the patient during procedures.	Requires Spline Assy, A Size (set of 2) mounted to the Back Spline Location.
Foot Control, 4 Function (Leg, Back, Lift, Tilt)	Provides a control device operated by foot.	

LIST OF ACCESSORIES

DESCRIPTION	USE	RESTRICTIONS
Foot Control Slant Mount Assy, Mobile Base	Provides a slanted surface to mount a foot control at the back of the stretcher bed.	
Foot Mounted Steering Handles	Provides (2) handles mounted at the foot of the stretcher bed for positioning or transport	
Forearm Support f/Deep Oval Headrest Assy (set of 2)	Provides a forearm support during chinning procedures.	Requires Deep Oval Headrest.
Forearm Support f/Prosthetic Headrest Assy (set of 2)	Provides a forearm support during chinning procedures.	Requires Prosthetic Headrest.
Head Stabilizer Cushion, Ophthalmic Headrest Mtd	Supports/stabilizes patient's head during procedure.	Requires Ophthalmic Headrest.
IV Rod Assy, Adj	Provides an adjustable IV rod.	Requires Accessory Mount, Base.
IV Rod Assy, Adj, Post Mtd	Provides an adjustable IV rod mounted to the post on a base mounted accessory.	Requires Mayo Tray Assy, Mayo Tray with Light Adptr Assy, or Cardiac Monitor Tray Assy.
Mayo Tray Assy, Base Mount Mtd	Provides a tray for the placement of small instruments and other items.	Requires Accessory Mount, Base. Not compatible on the same side as a Mtg Bracket Assy or Electrosurgery Generator.
Mayo Tray w/Light Adptr Assy, Base Mount Mtd	Provides a tray for the placement of small instruments and other items and a mount for an Exam Light, Lumerus LED, Flex/Rigid Neck	Requires Accessory Mount, Base. Not compatible on the same side as a Mtg Bracket Assy, Electrosurgery Generator.
Monitor Mount Assy w/Post, Base Mount, Cardiac	Mounts a patient monitor for viewing during procedures.	Requires Accessory Mount, Base.
Mtg Bracket Assy, Electrosurgery Generator	Provides a bracket to mount an Electrosurgery Generator to the base of a stretcher bed.	Not compatible on the same side as an Accessory Mount, Base.
Oxygen Tank Holder	Provides a support for mounting an Oxygen Tank to the rear of the stretcher bed.	Only compatible with 4.3" diameter tanks
Oxygen Tank Holder f/Foot Control Slant Mount	Provides a support for mounting an Oxygen Tank to the rear of the stretcher bed and attached to the foot control slant mount.	Only compatible with 4.3" diameter tanks. Requires Foot Control Slant Mount
Paper Roll Holder Assy, Adjustable 18" or 21"	Used for mounting a paper roll to the back of the stretcher bed.	Not compatible with a back mounted Exam Light.
Post Assy w/Light Adptr, Base Mount	Provides a post for mounting an Exam Light, Lumerus LED, Flex/Rigid Neck, 100-240V.	Requires an Accessory Mount, Base.
Power Cord, IEC w/Hospital Grade 90 deg Plug, 3 ft	Provides a 3' power cord with a 90 degree plug.	
Spline Assy, A Size (set of 2)	Provides 2 splines to mount in various places on the stretcher bed.	Not used on the Rear Seat Spline Location.
Spline Assy, B Size (set of 2)	Provides 2 splines to mount in the Rear Seat location.	Only used in the Rear Seat Spline Location.
Stirrups, Seat Spline Mtd (pair)	Used to position patient's legs during procedures.	Requires Spline Assy, A Size (set of 2) mounted in Front Seat Spline Location.

LIST OF ACCESSORIES

DESCRIPTION	USE	RESTRICTIONS
Strap, Body Restraint	Provides a strap to secure the patient's body.	
Strap, Leg Restraint	Provides a strap to secure the patient's legs.	
Transfer Support, ADA Compliant	Used as a support to assist the patient in transferring to the transfer surface.	Cannot be inserted on the same side of the chair while any accessory is mounted in the floating arm spline. Cannot be inserted on the same side of the chair as any seat rail mounted accessory or a Bed Rail & Transfer Support, ADA Compliant.
Vision Block, Seat Spline Mtd	A curtain to shield procedures from the patient's view.	Requires Spline Assy, A Size (set of 2) mounted in Front Seat Spline Location.
Wrist Support Assy, Crescent	Provides an adjustable support for wrists during procedures.	Requires Ophthalmic Headrest.
Wrist Support Assy, U-Shaped	Provides an adjustable support for wrists during procedures.	Requires Ophthalmic Headrest.

MAINTENANCE

⚠ WARNING/AVERTISSEMENT

There are no user serviceable parts inside. Refer all servicing to a qualified service technician.

Aucune pièce interne ne peut être réparée par l'utilisateur. Adressez-vous à un technicien qualifié.

⚠ WARNING/AVERTISSEMENT

After equipment is serviced, verify programmable positions (Home/P1-P4) are safe and known positions.

Après l'équipement est installé, vérifier les positions programmables (position retournée/P1-P4) sont des positions sûres et connues.

⚠ WARNING/AVERTISSEMENT

The Control of Hazardous Energy (lockout/tagout) 29 CFR §1910.147 procedure must be followed to disable equipment, preventing the release of hazardous energy while performing servicing and maintenance activities.

La procédure de La Maîtrise des Énergies Dangereuses (lockout/déconsignation) 29 CFR §1910.147 doit être suivie pour désactiver l'équipement, empêchant la libération des énergies dangereuses lors de l'exécution des activités de maintenance et d'entretien.

⚠ WARNING/AVERTISSEMENT

MTI recommends cleaning the equipment frequently. Follow the current CDC Guideline for Disinfection & Sterilization in Healthcare Facilities or other local guidelines for disinfecting.

MTI recommande de nettoyer l'appareil fréquemment. Suivez les lignes directrices actuelles du CDC pour la désinfection et la stérilisation dans les établissements de santé ou d'autres directives locales pour la désinfection.

⚠ CAUTION/PRÉCAUTION

This equipment may be damaged by solvents and dyes. Immediately remove any fluids spilled on the equipment.

Cet équipement peut être endommagé par des solvants et des colorants. Retirer immédiatement tout liquide renversé sur l'équipement.

⚠ CAUTION/PRÉCAUTION

Contact with sharp or pointed objects can damage the upholstery. Do not place hard surface objects on the upholstery surfaces as temporary indentations or impressions in the upholstery cover will result. Prolonged contact with a hard or irregular shaped object may make the impression semi-permanent.

Le contact avec des objets pointus ou tranchants peut endommager le capitonnage. Ne placez pas d'objets de surfaces dures sur les surfaces de capitonnage d'ameublement, comme des empreintes temporaires ou des impressions dans le couvercle du capitonnage en résulteront. Un contact prolongé avec un objet en forme de disque ou irrégulière peut faire l'impression semi-permanente.

⚠ CAUTION/PRÉCAUTION

Any sprayed disinfectant or cleaning solution should be wiped up immediately after application and not let to stand. A small sample area should be tested before any large application of any cleaner/disinfectant.

Tout désinfectant pulvérisé ou solution de nettoyage doivent être essuyés immédiatement après l'application et ne pas laisser reposer. Une petite zone de l'échantillon doit être testé avant toute application complète d'un nettoyant/désinfectant.

⚠ CAUTION/PRÉCAUTION

Avoid prolonged direct sunlight exposure to preclude decolorization.

Évitez l'exposition prolongée au soleil direct afin d'éviter la décoloration.

⚠ CAUTION/PRÉCAUTION

Replace damaged upholstery to prevent cross contamination.

Remplacer le capitonnage endommagé pour éviter la contamination croisée.

Disinfection

Follow the current CDC Guideline for Disinfection & Sterilization in Healthcare Facilities or other local guidelines.

Upholstery

MTI understands that many customers prefer to use a pre-mixed disinfecting solution in a spray or wipe. It is MTI's recommendation not to use sprays because over time the overspray will cause damage to the surfaces if not immediately wiped up. See "MTI Suggested Disinfectants Policy" for more details on disinfecting solutions which have been tested on MTI's products. It is available for download at www.mti.net/DisinfectantsPolicy. In the U.S. & Canada, call

(877) 875-4990, or International call (801) 875-4990, or email technical.support@mti.net to request a copy.

Cleaning

Upholstery should be cleaned weekly with a mild liquid soap and water mixture. Afterwards, wipe upholstery with clean damp cloth.

Painted and Plastic Surfaces

All painted metal parts have been electrostatically painted and the paint is baked on for durability. Weekly, those surfaces should be wiped down with a quality grade cleaner and a soft cloth or a damp cloth with mild soap.

Preventative Maintenance

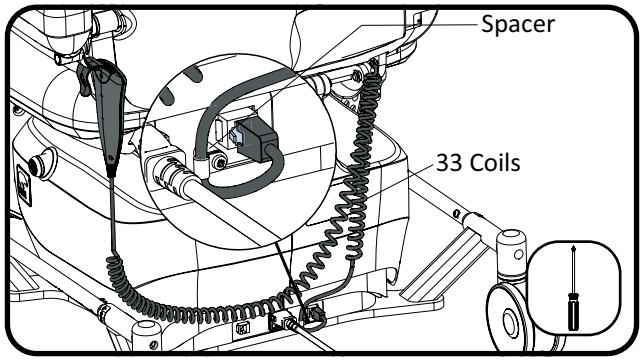
Periodically inspect the following areas:

1. All fasteners and pins should be in place and tightened securely.
2. Power cord should be free of cuts or other visible damage.
3. All mechanical functions should operate properly.

Have an authorized service technician inspect your equipment once a year.

Hand Control Cord

1. The hand control cord comes pre-attached to the back and base of the stretcher bed to prevent the cord from being damaged. If the hand control is removed, the hand control should be reattached with 33 coils between the clamps to prevent damage. The lower clamp should be attached with the spacer between it and the base.



Power Cord Removal Instructions

1. To remove the power cord, pull back on the red tabs of the power cord at the base of the stretcher bed. See “Unpacking and Set-up” on page 12 for installation instructions.

Fuse Replacement Instructions

1. See Figure 3. Pull back on the red tabs of the power cord at the base of the stretcher bed to remove the power cord.
2. Using a flat head screwdriver, open the fuse drawer in the power cord jack.
3. Remove the burnt fuse(s).
4. Insert new fuse(s) into the fuse drawer with same size fuse (5mm x 20mm 4 Amp 250V Time-Delay Glass tube type) removed in step 3 (See Figure 3).
5. Reinsert the fuse drawer. Ensure it is completely inserted.
6. Reinsert the power cord into the base of the stretcher bed.

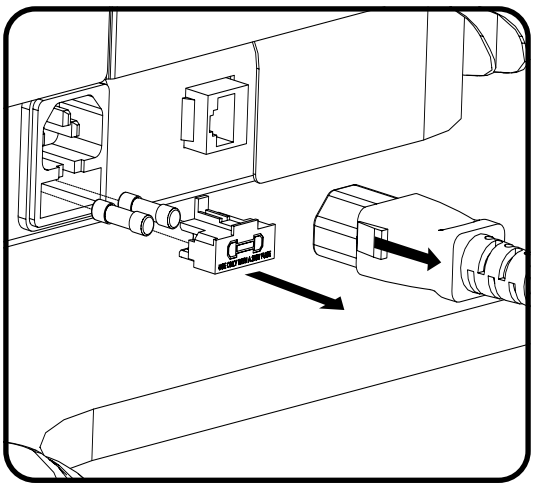


Figure 3

Calls for Service

Equipment’s model and serial number information are required when calling for service.

Serial number is indicated by the SN symbol located on the serial number tag.

If service is required, contact MTI.

U.S. & Canada (877) 875-4990
International (801) 875-4990
Fax (801) 952-0548
technical.support@mti.net
www.mti.net

LIMITED WARRANTY

Medical Technology Industries, Inc. (hereinafter referred to as MTI), shall repair or replace products of its manufacture, which prove to be defective in material and/or workmanship for a period of twelve (12) months from the date of shipment to the customer (the "Warranty Period"). MTI will, at its option, provide parts to the customer or repair the defective part at MTI's factory or authorized repair facility. Except as it may otherwise specifically agree in writing, MTI shall not be liable for transportation or labor charges for repairs or adjustments or other work done on any of its products by MTI's authorized dealers or service organizations.

Customer shall notify MTI of any alleged claim or defect within fourteen (14) days from the date customer discovers, or upon reasonable inspection should have discovered, such alleged claim or defect (but in any event before the expiration of the Warranty Period). All repair parts are warranted for Ninety (90) calendar days from the date of shipment to the customer. Warranty is non-cumulative.

This warranty does not apply to any part or product which upon examination by MTI or its authorized agents indicates the product has been repaired or altered in any way by persons other than MTI or its authorized agents, has been misused, abused or used in a manner contrary to any instructions issued by MTI. Any parts or products claimed under this warranty must be properly packaged by customer and returned to MTI's factory with no liability to MTI for the parts, products or transportation charges thereon.

In no event, shall MTI have any liability for consequential, indirect, incidental, special, exemplary, punitive or enhanced damages, lost profits or diminution in value arising out of or relating to a product defect or the repair or replacement of any products manufactured or sold by it.

THE FOREGOING LIMITED WARRANTY IS MADE IN LIEU OF ALL WARRANTIES EXPRESSED OR IMPLIED, AND MTI MAKES NO OTHER WARRANTY WHATSOEVER WITH RESPECT TO THE PRODUCTS INCLUDING BUT NOT LIMITED TO, ANY IMPLIED WARRANTY OR MERCHANTABILITY OR FITNESS FOR A PARTICULAR PURPOSE.

TROUBLESHOOTING GUIDE

NOTICE/AVIS

See "GUIDE TO INDICATOR LIGHTS AND SOUNDS" on page 31 for additional solutions. For more information on this equipment, including circuit diagrams, parts and service manuals (Item No. 130-3251), and other helpful information, please contact MTI.

Voir «GUIDE TO INDICATOR LIGHTS AND SOUNDS» à la page 31 pour des solutions supplémentaires. Pour plus d'informations sur cet équipement, y compris des schémas de circuits, les manuels de service et pièces (Numéro d'article 130-3251), et d'autres information utiles, s'il vous plaît contacter MTI.

#	SYMPTOMS	PROBLEM CAUSE	SOLUTION
1	Stretcher bed will not function on AC Power (works on battery power). Hand control lights are off. No green light visible on power supply through back of stretcher bed.	Stretcher bed power cord is not securely connected.	Verify power cord is securely connected at both ends.
		Facility power or receptacle problem. Facility power circuit breaker problem.	Check facility's power receptacle with a known working device of same voltage specifications. Also check facility circuit breaker to verify it is not tripped if no power to known working device.
		Facility supply voltage is incorrect voltage.	Verify voltage is per specifications on stretcher bed serial number/product tag.
		Fuses have blown.	Verify fuses are not burnt. Replace fuse(s) if necessary. See "Fuse Replacement Instructions" on page 27.
		Stretcher bed Power Supply is faulty.	Contact Technical Support.
2	Stretcher bed will not function on battery power (works on AC power). Hand control lights are off. No green light visible on power supply through back of stretcher bed.	Battery is not fully inserted.	Confirm the battery is fully inserted.
		Battery is low or out of power.	Check battery fuel gauge. Swap or charge battery if necessary.
3	Stretcher bed will not function. Hand control lights are off. Green light visible on power supply through back of stretcher bed.	Stretcher bed is off.	Press the "On" Button on the hand control or unplug the stretcher bed for 30 seconds and plug it back in.
		Faulty Hand Control	Test an alternate hand control from similar equipment. Test foot control from similar equipment.
		Faulty Hand Control Jack	Test the same hand control in the foot control jack.
		Stretcher bed PCA isn't functioning correctly.	Contact Technical Support.
4	The stretcher bed suddenly loses power while the stretcher bed is operating on battery power.	Loads on the stretcher bed have caused the battery to shut off.	Battery will reset after approximately 1 minute.
5	Lift, Back, Tilt, or Foot functions will not function.	An actuator is experiencing excessive loads.	Remove or reposition patient, move the stretcher bed in opposite direction, then in the intended direction to continue normal operation.
		An actuator is not functioning or Stretcher bed PCA isn't functioning correctly.	Contact Technical Support.

TROUBLESHOOTING GUIDE

NOTICE/AVIS

See "GUIDE TO INDICATOR LIGHTS AND SOUNDS" on page 31 for additional solutions. For more information on this equipment, including circuit diagrams, parts and service manuals (Item No. 130-3251), and other helpful information, please contact MTI.

Voir «GUIDE TO INDICATOR LIGHTS AND SOUNDS» à la page 31 pour des solutions supplémentaires. Pour plus d'informations sur cet équipement, y compris des schémas de circuits, les manuels de service et pièces (Numéro d'article 130-3251), et d'autres information utiles, s'il vous plaît contacter MTI.

#	SYMPTOMS	PROBLEM CAUSE	SOLUTION
6	Foot Extension will not extend and beeps.	The footrest is at the safe zone boundary. The footrest is too close to the floor or is too close to vertical.	Change the angle of the footrest.
7	The stretcher bed consistently beeps when retracting the foot section.	The footrest is at the safe zone boundary. The footrest is avoiding colliding with the seat cushion.	This is normal operation.
8	The stretcher bed beeps when pressing Lift Down or Tilt Down and stops moving.	The footrest is too close to the floor.	Press the Foot Retract or Foot Up Button to move it away from the floor.
9	The stretcher bed beeps when pressing Back Down or Tilt Up and stops moving.	The backrest has reached its safe zone boundary. It is avoiding potential collisions.	Press the Lift Up Button to raise the top of the stretcher bed above the safe zone boundary or alter the position of the back or tilt.
10	The stretcher bed will not function. The green light and orange light are lit on the hand control or stretcher bed PCA.	The Safety Lockout Feature is enabled.	To exit Safety Lockout press **Stop, Stop, Back Down, Stop.**
11	The stretcher bed does not move when a Program Button is pressed.	The Program Button Setting is set to the Maintain Button Setting.	See "Smart Controller™ Technology" on page 13 for instructions to change the Program Button Setting.
		The Program Button does not have a saved location.	See "Controls" on page 15 for instructions on saving a Program Button location.
12	Home Button does not move the stretcher bed to the expected location.	The saved location for the Home Button has been set by another user/ inadvertently.	Move the stretcher bed to the desired location. See "Controls" on page 15 for instructions on saving a Home Button location. For further instructions about the operation of the button, see "Smart Controller™ Technology" on page 13.
13	The stretcher bed consistently beeps while moving to a programmed position.	An actuator is malfunctioning.	Contact Technical Support.
14	Stretcher bed will not function or a function is erratic.	Stretcher bed PCA isn't functioning correctly.	Contact Technical Support.
15	Stretcher bed PCA relays turn on/off quickly and excessively ("chatters").	Stretcher bed PCA or hand/foot control may be functioning incorrectly.	Contact Technical Support.
16	Lift function is slower than usual and only goes up about halfway. Stretcher bed beeps while moving.	One of the lifting column motors is not functioning or stretcher bed PCA isn't functioning correctly.	Contact Technical Support.
17	Motor is running after stretcher bed stops.	A motor gear has stripped.	Contact Technical Support for motor replacement.
18	The stretcher bed will roll but not steer.	Both locking bars are in STEER position on a stretcher bed with Selective Steer Option.	Change one of the locking bars to the FREE position. See page 18 for instructions.

GUIDE TO INDICATOR LIGHTS AND SOUNDS

#	LIGHTS						SOUND	STATUS OR CONDITION
	GREEN		BLUE		ORANGE			
1	●	Solid	○	Off	○	Off	Start Up Tune	Stretcher bed has been plugged in.
2	✸	Slow Flash	○	Off	○	Off	None	Stretcher bed is in hibernation state to conserve power. To exit hibernation press any button.
3	●	Solid	✸	Rapid Flash	○	Off	None	Stretcher bed is moving to the Programmed Home or a Program Position or manual button (Lift Up/Down, Tilt Up/Down, Back Up/Down, Foot Up/Down, or Foot In/Out) is being pressed.
4	●	Solid	✸	Rapid Flash	✸	Rapid Flash	None	Stretcher bed is moving to the ADA Compliant Home position.
5	●	Solid	○	Off	○	Off	High Beep when a Program Button is pressed	No Program Position is saved.
6	●	Solid	✸	Flash Twice	○	Off	High-Low-High Tune	Program Position or Programmed Home Position has been saved.
7	●	Solid	○	Off	✸	Mod. Flash for 10 sec.	High Beep during motion of specific actuator	Actuator/motor problem (Contact Technical Support).
8	●	Solid	○	Off	✸	Flash Twice	High Beep	Stretcher bed has been moved to a Safe Zone boundary. (ex. Excessive Trendelenburg or Footrest near Seat Cushion or floor). Move in opposite direction to resume full functionality.
9	●	Solid	○	Off	✸	Flash Twice	High Beep	Excessive load on a motor. Remove or shift excess load and move in the opposite direction momentarily to resume.
								Actuator/motor reaches end of stroke prematurely. NOTE: If beeping occurs frequently, contact Technical Support.
10	●	Solid	○	Off	✸	Rapid Flash	None	Swivel has been unlocked. NOTE: on this non-swivel stretcher bed model, the orange LED will still flash when the Swivel Button is pressed. Press any button to turn it off.
11	●	Solid	○	Off	●	Solid	2 High Beeps (upon activation)	Safety Lockout feature is enabled. (To exit press Stop, Stop, Back Down, Stop) Orange LED will flash twice each time button is pressed while Safety Lockout feature is enabled.
12	●	Solid	○	Off	✸	Slow Flash	High Beep with every button press	No actuators respond to motion commands. Chair ID Mismatch (Contact Technical Support).
13	✸	Mod. Flash	✸	Mod. Flash	✸	Mod. Flash	High Beep with every button press	No actuators respond to motion commands. Chair ID Cable Invalid (Contact Technical Support).
14	○	Off	○	Off	○	Off	None	Loss of power (stretcher bed was unplugged and/or battery was removed).

GUIDE TO INDICATOR LIGHTS AND SOUNDS

#	LIGHTS						SOUND	STATUS OR CONDITION
	GREEN		BLUE		ORANGE			
15	○	Off	○	Off	○	Off	Shut Down Tune	Stretcher bed has been powered off. Press the On/Off Button on hand control to resume operation.
16	●	Solid	✸	Flash Twice	✸	Flash Twice	Low Beep	Stop Button has been pressed for 5 seconds awaiting sequence to enter Smart Controller™ User Program Mode or Calibration Mode.
17	●	Solid	✸	Flash once	○	Off	High Beep	High Beep with each button press until sequence has been entered for Smart Controller™ User Program Mode or Calibration Mode.
USER PROGRAM MODE								
18	●	Solid	●	Solid	✸	Slow Flash	Low-High Tune then Low Beep every 10 sec.	Stretcher bed has entered Smart Controller™ User Program Mode. Exits automatically after 30 seconds of no activity.
19	●	Solid	○	Off	✸	Slow Flash	High Beep each button press	Button press while in Smart Controller™ User Program Mode.
20	●	Solid	✸	Flash Twice	✸	Slow Flash	High-Low-High Tune	New setting accepted.
21	●	Solid	●	Solid	✸	Slow Flash	Three Low Beeps	New setting not accepted.
22	●	Solid	○	Off	○	Off	High-Low Tune	Stretcher bed has exited Smart Controller™ User Program Mode.
CALIBRATION MODE								
23	○	Off	○	Off	✸	Mod. Flash	High Beep when an individual motor button is pressed.	One or more actuators is out of calibration. Do not operate the stretcher bed. Contact Technical Support.
24	✸	Slow Flash	✸	Slow Flash	✸	Slow Flash	Low-High Tune then Low Beep every 10 sec.	Stretcher bed is in Calibration Mode. Do not operate the stretcher bed. Contact Technical Support.

SPECIFICATIONS

DESCRIPTION	SPECIFICATION
Load Capacity[1]	
Lift	725 lb (328.9 kg)
Back	400 lb (181.4 kg)
Tilt	725 lb (328.9 kg)
Foot (footrest fully extended)	150 lb (68 kg)
Foot (footrest retracted)	200 lb (90.7 kg)
Weight of stretcher bed[2]	294 lb (133.4 kg)
Shipping Weight of stretcher bed[3]	361 lb (163.8 kg)
Shipping Weight of Upholstery Kit	
840 Seamless/Designer Upholstery Kit	28 lb (12.7 kg)
Table Top Range	18.75-37 in (47.6-94 cm)
Back Section Range	0°± 2.2° (horizontal) to 85°± 2.2°
Table Top Tilt Range	0° ± .6° to 30°± .6° Range is limited depending on Back angle and Lift position.
Foot Section Range	0° ± 2.2° (horizontal) to 90°± 2.2°
Foot Extension Range	7 in (17.78 cm) Range is limited depending on Tilt angle, Lift, and footrest angle.
Foot Control	IPX6 Rated
Hand Control	IPX6 Rated
General Dimensions (w/headrest & footrest fully extended	25 in (63.5 cm) wide x 80 in (203.2 cm) long
Power Cord	Plugs: U.S.A. standard 3 prong and IEC 60320-C13 (self-retaining)
Electrical Ratings	100-240VAC (47-63 Hz); 7 Amps
Battery Specifications	Charging Temperature Range: 0° C to 45° C Voltage: 28.8V ● Max Current: 12 Amps ● Capacity: 2.5 Ah
Fuse Replacement	(2) 5mm x 20mm 4 Amp 250V Time-Delay Glass Tube Fuse
Duty Cycle	Intermittent Operation, 10% Duty Cycle Example: (1) minute on (9) minutes off
Classification	Class 1, Type B Applied Part, Ordinary Equipment, Intermittent Operation
Listing/Approvals	ETL, IEC 60601-1:2005 Ed.3+A1;C1:2014 AAMI ES60601-1:2005+A1 CSA C22.2#60601-1:2014 Ed.3 IEC 60601-1-2:2014 Ed.4

[1]Load Capacity will vary depending on placement and distribution of load on the section of the stretcher bed.

[2]Weight may vary due to product variation.

[3]Weight may vary due to variation in weight of shipping materials.

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In-service Checklist for G2 Products with SmartTech™

Model Number	Description	Serial Number(s)

The following in-service needs to be performed on this MTI product:

Set Smart Controller™ Settings

Enter the User Program Mode in the Smart Controller™ to change how the Program Buttons operate. See "User Program Mode" on page 14.

Button Setting Momentary Maintain

[Factory default] *Momentary Button Setting: press and release to automatically move bed to the programmed position. Maintain Button Setting: press and hold to move bed to the programmed position. Once button is released, bed will stop.*

Reviewed Battery Use and Charging

2 Batteries are rotated between use. Charge depleted battery at the end of day. Insert the other battery at start of day. Remove battery from bed when not in use to increase use time. Review "Battery & Charger Operation" on page 20.

Saved Programmable Home

Save by positioning bed to desired location. If buttons are set to Momentary Button Setting, press and hold the Home Button for 10 seconds. If buttons are set to Maintain Button Setting, press the Stop Button twice and then press and hold the Home Button for 10 seconds. Once saved, audible and visual confirmation will be given.

Saved (4) User Program Buttons (P1-P4)

Save by positioning bed to desired location. If buttons are set to Momentary Button Setting, press and hold the desired program button for 5 seconds. If buttons are set to Maintain Button Setting, press the Stop Button twice then press and hold the desired program button for 5 seconds. Once saved, audible and visual confirmation will be given.

Review Mobile Base Operation

Identify the Mobile Base Option - Front Steer, Rear Steer, or Select Steer
Review Pedal Settings for Steering Mode, Free Mode, and Lock Mode.

Reviewed ADA Compliant Home

ADA Compliant Home position is Lift Down, Tilt Down, Back Up, and Foot Down and Retracted. To access: if buttons are set to Momentary, press and release the Home Button twice (within 1 second). If buttons are set to Maintain, press and hold the Home Button to move to Programmable Home Position, then release and press and hold the Home Button again (within 1 second) until the stretcher bed stops moving.

Reviewed Safety Lockout

Locks out all commands once this state has been enabled. To enter, press the following sequence of buttons: Stop, Stop, Back Up, Stop. Once enabled, audible and visual confirmation will be given.
To exit, press the following sequence of buttons: Stop, Stop, Back Down, Stop. Once disabled, audible and visual confirmation will be given.

Reviewed Controls and Operation sections of the User Manual

If you would like to receive occasional emails regarding MTI promotions and product announcements, fill out the sales form at www.mti.net/contact-us.

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