Recommended separation distances between portable and mobile RF communications equipment and this device:

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

<table>
<thead>
<tr>
<th>Rated maximum output power of transmitter (W)</th>
<th>Separation distance according to frequency of transmitter (m)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>150 kHz to 80 MHz</td>
</tr>
<tr>
<td>0.01 W</td>
<td>$d = 1.2 \sqrt{P}$</td>
</tr>
<tr>
<td>0.1 W</td>
<td>0.12</td>
</tr>
<tr>
<td>1 W</td>
<td>0.38</td>
</tr>
<tr>
<td>10 W</td>
<td>1.2</td>
</tr>
<tr>
<td>100 W</td>
<td>3.8</td>
</tr>
<tr>
<td>1200 W</td>
<td>12</td>
</tr>
</tbody>
</table>

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:**

This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it is used in such an environment.

<table>
<thead>
<tr>
<th>Immunity Test</th>
<th>IEC60601 test level</th>
<th>Compliance</th>
<th>Electromagnetic Environment-Guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Conducted RF</td>
<td>IEC 61000-4-6</td>
<td></td>
<td>Portable and mobile RF communications equipment should be used no closer to any part of this device, including cables, than there commended separation distance calculated from the equation applicable to the frequency of the transmitter.</td>
</tr>
<tr>
<td>Radiated RF</td>
<td>IEC 61000-4-3</td>
<td></td>
<td>Recommended separation distance $d = \frac{1.2P}{150\text{kHz} \text{ to } 80\text{MHz}}$</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>$d = \frac{1.2\sqrt{P}}{150\text{kHz} \text{ to } 80\text{MHz}}$</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>$d = \frac{2.3\sqrt{P}}{80\text{MHz} \text{ to } 2.5\text{GHz}}$</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td>Where $P$ is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and $d$ is the recommended separation distance in meters (m).</td>
</tr>
</tbody>
</table>

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.

### IMPORTANT SAFEGUARDS

**READ ALL INSTRUCTIONS BEFORE USE**

**DANGER - To reduce the risk of electrical shock:**
1. Always unplug this product immediately after using.
2. Do not use while bathing.
3. Do not place or store product where it can fall or be pulled into a tub or sink.
4. Do not place in or drop into water or other liquids.
5. Do not reach for a product that has fallen into water. Unplug immediately.

**WARNING - To reduce the risk of burns, electrical shock, fire, or injury:**
1. This product should never be left unattended when plugged in.
2. Close supervision is necessary when this product is used by, on, or near children or invalids.
3. Use this product only for its intended use as described in this manual.
4. Do not use attachments not recommended by the manufacturer.
5. Never operate this product if it has a damaged cord or plug, if it is not working properly, if it has been dropped or damaged, or exposed to water. Return the product to a service center for examination and repair.
6. Keep the cord away from heated surfaces.
7. Never block the air openings of this product. Never place the product on a soft surface, such as a bed or couch, where the air openings may become blocked. Keep the air openings free of lint, hair, and other similar particles.
8. Never drop or insert any object into any opening or hose on this product.
9. Connect this product to a properly grounded outlet only. See Grounding Instructions.

**NOTES, CAUTIONs AND WARNINGS:**

**NOTE -** Calls attention to useful information, tips, reminders.

**CAUTION -** Calls attention to correct operating or maintenance procedures in order to prevent damage to or destruction of the equipment or other property.

**WARNING -** Calls attention to a potential danger that requires correct operating procedures or practices in order to prevent personal injury.

**SYMBOL Definitions**

- **EC**
  - Authorized representative in the European community.

- **Manufacture**
  - Complies with standards protecting against electric shock for type BF equipment.

- **Attention, you should read the accompanying information carefully!**

- **Attention – Observe proper Disposal of Electrical & Electronic Equipment (WEEE):**
  - This product should be handed over to an appropriate collection point for the recycling of electrical and electronic equipment. For more detailed information about the recycling of this product, please contact your local city office, household waste disposal service or the retail store where you purchased this product.

- **Consult operating instructions**
  - Class II 

- **Functional earth**
1. INTRODUCTION
This manual should be used for initial set up of the system and for reference purposes.

1.1 GENERAL INFORMATION
This system is a high quality, affordable mattress system, suitable for the treatment and prevention of pressure ulcers. This product has been tested and successfully approved for the following standards:

EN 60601-1
EN 60601-1-2
EN 55011 Class B
IEC61000-3-2
IEC 61000-3-3

FOR US AND CANADA ONLY

E245395
53DG

Medical Equipment-Air Pump
with respect to electrical shock, fire and mechanical hazards only in accordance with UL60601-1 AND CAN/CSA C22.2 No.601.1

Le produit à été testé avec des équipements médicaux et respecte les normes UL 60601-1 & CAN/CSA C22.2 No.601.1, prévenant les choc électrique, le feu et les risques de blessures physiques.

1.2 INTENDED USE
This product is intended:
- to help and reduce the incidence of pressure ulcers while optimizing patient comfort.
- for long term home care of patients suffering from pressure ulcers.
- for pain management as prescribed by a physician.

NOTE: This equipment is not suitable for use in the presence of flammable anesthetic mixture with air or with pure oxygen or nitrous oxide.

NOTE: L’équipement ne peut être utilisé s’il y a risque de mélange d’un anesthésique inflammable avec l’air ou l’oxygène ou oxyde nitreux.

APPENDIX A: EMC INFORMATION

Guidance and Manufacturer’s Declaration- Electromagnetic Emissions:
This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it is used in such an environment.

<table>
<thead>
<tr>
<th>Emissions Test</th>
<th>Compliance</th>
<th>Electromagnetic Environment-Guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Harmonic emissions</td>
<td>Class A</td>
<td>The device is suitable for use in all establishments, including domestic ones and those directly connected to the public low-voltage power supply network.</td>
</tr>
<tr>
<td>IEC61000-3-2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Voltage fluctuations / Flicker emissions</td>
<td>Complies</td>
<td></td>
</tr>
<tr>
<td>IEC61000-3-3</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

Guidance and Manufacturer’s Declaration- Electromagnetic Immunity:
This device is intended for use in the electromagnetic environment specified below. The user of this device should make sure it is used in such an environment.

<table>
<thead>
<tr>
<th>Immunity Test</th>
<th>IEC60601 test level</th>
<th>Compliance</th>
<th>Electromagnetic Environment-Guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>Electrostatic Discharge (ESD)</td>
<td>±6kV contact</td>
<td>±6kV air</td>
<td>Floors should be wood, concrete or ceramic, the relative humidity should be at least 30%.</td>
</tr>
<tr>
<td>IEC61000-4-2</td>
<td>±6kV contact</td>
<td>±6kV air</td>
<td></td>
</tr>
<tr>
<td>Voltage dips, short interruptions and voltage variations on power supply input lines</td>
<td>±20V line(s) to line(s)</td>
<td>±20V line(s) to earth</td>
<td>Mains power quality should be that of a typical commercial or hospital environment.</td>
</tr>
<tr>
<td>IEC61000-4-11</td>
<td>±1 kV line(s) to line(s)</td>
<td>±2 kV line(s) to earth</td>
<td></td>
</tr>
<tr>
<td>Voltage dips, short interruptions and voltage variations on power supply input lines</td>
<td>±5 % Uₜ (&lt;95 % dip in Uₜ) for 0,5 cycle</td>
<td>±5 % Uₜ (&lt;95 % dip in Uₜ) for 0,5 cycle</td>
<td>Mains power quality should be that of a typical commercial or hospital environment.</td>
</tr>
<tr>
<td>IEC61000-4-11</td>
<td>40 % Uₜ (&lt;60 % dip in Uₜ) for 5 cycles</td>
<td>40 % Uₜ (&lt;60 % dip in Uₜ) for 5 cycles</td>
<td></td>
</tr>
<tr>
<td>Voltage dips, short interruptions and voltage variations on power supply input lines</td>
<td>70 % Uₜ (&lt;30 % dip in Uₜ) for 25 cycles</td>
<td>70 % Uₜ (&lt;30 % dip in Uₜ) for 25 cycles</td>
<td></td>
</tr>
<tr>
<td>IEC61000-4-11</td>
<td>&lt;5 % Uₜ (&lt;95 % dip in Uₜ) for 5 sec</td>
<td>&lt;5 % Uₜ (&lt;95 % dip in Uₜ) for 5 sec</td>
<td></td>
</tr>
<tr>
<td>Power frequency (50/60Hz) - magnetic field</td>
<td>3 A/m</td>
<td>3 A/m</td>
<td>Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.</td>
</tr>
<tr>
<td>IEC61000-4-8</td>
<td></td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

NOTE: Uₜ is the a.c. mains voltage prior to the application of the test level
This product should be grounded. In the event of an electrical short circuit grounding reduces the risk of electric shock by providing an escape wire for the electric current. This product is equipped with a cord having a grounding wire with a grounding plug. The plug must be plugged into an outlet that is properly installed and grounded.

**DANGER** - Improper use of the grounding plug can result in a risk of electric shock.

If repair or replacement of the cord or plug is necessary, do not connect the grounding wire to either flat blade terminal. The wire with insulation having an outer surface that is green with or without yellow stripes is the grounding wire.

**NOTE** - If the repair or replacement of the cord is necessary, please contact a qualified electrician or serviceman. To reduce the risk of electric shock, do not modify the cord or plug in any way.

Check with a qualified electrician or serviceman if the grounding instructions are not completely understood, or if in doubt as to whether the product is properly grounded.

### 2. PRODUCT DESCRIPTION

#### 2.1 PUMP AND MATTRESS SYSTEM

1. CPR Tags
2. Pump unit
3. Mattress unit
4. Quick Connector

#### 2.2 PUMP UNIT

**Front View**
1. Quick Connector Slot
2. Front Panel

**Rear View**
1. Power Switch
2. Power Cord
3. Bed Frame Hangers
3. INSTALLATION

Unpack the box and check the package contents for completeness.

Package Content List
- Mattress unit x 1 (may not be included if only pump unit is purchased)
- Pump Unit x 1
- User Manual x 1

Inspect equipment for damage which may have occurred during shipment. If there is damage, please contact your dealer immediately.

3.1 PUMP & MATTRESS INSTALLATION

1. Place the mattress or pad on top of a bed frame. Orient the mattress so that the foot-markers on the mattress are situated towards the end of the bed (where pump will be located).

2. Place the pump hangers over the bed railing at the foot of the bed. Support the pump from the bottom and turn the hangers outward (or inward) to secure the pump against the railing.

   If no bed railing is available, place the pump on a flat, sturdy surface or on the floor, beneath the bed.

3. Connect the Quick Connector on the air hose (from air mattress) to the pump unit Quick Connector Slot. When you hear a "click", the connection will be secured.

   NOTE: Check and ensure the air hoses are not kinked or tucked under the mattress.

8. TECHNICAL DESCRIPTION

Specifications:

<table>
<thead>
<tr>
<th>Item</th>
<th>Specification</th>
</tr>
</thead>
<tbody>
<tr>
<td>Power Supply (Note: See rating label on the product)</td>
<td>AC 100-120V 60 Hz, 0.16 A 14W (For 120V System)</td>
</tr>
<tr>
<td>Fuse Rating</td>
<td>T1A, 250V</td>
</tr>
<tr>
<td>Cycle time</td>
<td>Fixed, 8 min @ 60Hz</td>
</tr>
<tr>
<td>Dimension (L x W x H)</td>
<td>29 x 18.5 x 12.6 (cm) or 11.4” x 7.3” x 5.0”</td>
</tr>
<tr>
<td>Weight</td>
<td>2.2 Kg or 5 lb</td>
</tr>
<tr>
<td>Environment</td>
<td></td>
</tr>
<tr>
<td>Temperature</td>
<td>Operation: 10° C to 40° C (50° F to 104° F)</td>
</tr>
<tr>
<td></td>
<td>Storage: -15° C to 50° C (5° F to 122° F)</td>
</tr>
<tr>
<td></td>
<td>Shipping: -15° C to 70° C (5° F to 158° F)</td>
</tr>
<tr>
<td>Humidity</td>
<td>Operation: 10% to 90% non-condensing</td>
</tr>
<tr>
<td></td>
<td>Storage: 10% to 90% non-condensing</td>
</tr>
<tr>
<td></td>
<td>Shipping: 10% to 90% non-condensing</td>
</tr>
<tr>
<td>Classification</td>
<td>Class II with functional earth(NOTE: 5*), Type BF, IPX0</td>
</tr>
<tr>
<td></td>
<td>Not suitable for use in the presence of a flammable anesthetic mixture (No AP or APG protection)</td>
</tr>
<tr>
<td>Mattress</td>
<td></td>
</tr>
<tr>
<td>Model</td>
<td>8” Replacement</td>
</tr>
<tr>
<td>Dimension (L x W x H)</td>
<td>200 x 90 x 20.3 (cm) or 78.7” x 35.4” x 8”</td>
</tr>
<tr>
<td>Weight</td>
<td>7.5 kg 16.5 lb</td>
</tr>
<tr>
<td>Pressure Range</td>
<td>Adjustable Weight Settings</td>
</tr>
</tbody>
</table>

NOTE: 1. Consult the distributor or EU representative for further technical documents.
2. These specifications are also applicable for other regions operating with the same power supply.
3. The manufacturer reserves the right to modify the specifications without notice.
4. The earthing terminal is served as functional earth, not protective earth.
6. MATTRESS STORAGE

1. Lay the mattress on a flat surface, upside-down.
2. Roll-up the mattress starting from the head until you reach the foot of the mattress.
3. Use the straps at the end of the mattress and stretch them around the mattress to prevent it from unraveling.

⚠️ Note: Do not fold, crease or stack the mattresses.

7. MAINTENANCE

7.1 GENERAL

1. Check the main power cord and plug for abrasions or excess wear.
2. Check the mattress cover for signs of wear or damage.
3. Disconnect the air tube from the mattress. Then check the airflow coming from the two air outlets on the pump. They should be alternately delivering air when the pump is set at “alternate” mode.
4. Check the air hoses for kinks or breaks. For replacement, please contact your local distributor.

4. Plug the power cord into a electrical outlet with the appropriate voltage.

⚠️ NOTE: The pump unit must be suitable for the local power voltage.

⚠️ NOTE: S’assure que la pompe est compatible au voltage local ou disponible.

6. Turn the main power switch (on pump) to the ON position. After you turn on power to the pump, you will have to wait at least 20 minutes for the mattress to become fully inflated.

The low pressure indicator (yellow LED) will illuminate while the mattress is pressurizing.

When the appropriate pressure is reached, the Low Pressure indicator (yellow LED) will go off and the Normal Pressure indicator (green LED) will illuminate. The bed is now ready for use.

⚠️ NOTE: You can unplug the unit to power-off the device.

⚠️ NOTE: L’appareil est également muni d’une fiche de connexion à l’électricité.

⚠️ CAUTION: The pump supplied with the mattress can only be used for mattresses recommended by the manufacturer. Do not use it for any other purpose. (applied part: air mattress)

⚠️ AVERTISSEMENT : La pompe ne peut être utilisée qu’avec le matelas designé ou recommandé par le manufactureur, Ne pas servier d’usage autre que celui auquel elle a été conçu

⚠️ NOTE: (For models WITHOUT the low-air-loss function) During power outages, you can cover the Quick Connector with the transport cap to maintain air pressure inside the air cells.

⚠️ NOTE: After installation, make sure any excess cord or tubing is removed from possible foot traffic to avoid accidental tripping. All EQUIPMENT should be positioned to always allow unhindered patient access by physicians and caregivers.
4. OPERATIONS

1. Pressure Adjustment Knob
   The pressure adjustment knob controls the air pressure in the mattress. Turning the knob clockwise will increase the pressure; counter-clockwise decreases the pressure. Higher pressures will support heavier patients. The pressure should be adjusted according to individual comfort preferences.

When the mattress is finished pressurizing to the desired level of firmness, the Normal Pressure LED indicator will illuminate.

NOTE: You can also check if the pressure is suitable for the patient by sliding one hand beneath the air cells at the level of the patient's buttocks. The air cells will alternately inflate and deflate. You should feel slight contact with the buttocks when the air cells beneath the buttocks deflate.

NOTE: Every time the mattress is initialized (inflated), it is recommended that you set the pressure knob to "Max" to hasten inflation. You can then adjust the air mattress to the desired firmness later.

2. Normal Pressure Indicator
   When the green LED illuminates, the pressure inside of air mattress has reached the desired pressure setting (preset on the Pressure Adjustment Knob).

3. Low Pressure Indicator
   When the mattress is initializing, the Low Pressure LED will illuminate until the appropriate pressure is reached (according to the Pressure Adjustment Knob); this is normal. Otherwise, the Low Pressure LED is a warning, indicating that the pressure in mattress is unusually low. Check to ensure all connections are secured and correctly installed as per the installation instructions.

4. Alternate / Static Switch
   The ALTERNATE/STATIC switch selects between Alternate Pressure mode and Static Pressure mode. With Alternate Pressure mode, alternating air cells are partially deflated and inflated, avoiding prolonged pressure on any single point beneath the patient; this is to prevent pressure ulcers. With Static Pressure mode, all of the air cells are equally inflated.

4.2 Emergency CPR Operations
   CPR must be performed on a firm surface. Therefore, if an emergency CPR situation occurs with the patient on the mattress, the mattress must be quickly deflated. This is done by quickly pulling on the CPR tag located at the head of the mattress on the patient's right-side. The quick connector found from the pump unit can also be disconnected to hasten deflation.

5. CLEANING
   It is important to follow the cleaning procedures before first-time use with any patient; this is to avoid the possibility of transmission of pathogens between patients and caregivers.
   Wipe down the pump unit with a damp cloth pre-soaked with a mild detergent. Avoid contact with dust and proximity to dusty areas. Make sure that any cleaning agents you use will not harm or corrode the plastic casing on the pump unit.

   CAUTION: Do not immerse or soak the pump unit in liquids.

   Wipe-down the mattress unit with a damp cloth pre-soaked with warm water (do not exceed 65 °C) containing a mild detergent. Avoid dust and proximity to dusty areas. The cover may also be cleaned using sodium hypochlorite diluted in water. All components should be air dried thoroughly before use.

   CAUTION: Do not use phenolic based products for cleaning.

   CAUTION: Do no dry the mattress in direct sunlight.

   The carrying bag (if available) should be turned inside-out and completely wiped-down using disinfectant solutions. Allow it to air-dry thoroughly and then turn it back and wipe down the outside of the bag with disinfectant solutions.