



Protekt® Aire 7000 Air Mattress System
Lateral Rotation, Alternating Pressure, Pulsation and
Low Air Loss.

User Manual



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IMPORTANT SAFEGUARDS

READ ALL INSTRUCTIONS BEFORE OPERATING THIS DEVICE



NOTE, CAUTION AND WARNING STATEMENTS:

NOTE – Indicate some tips.

CAUTION – Indicate correct operating or maintenance procedures in order to prevent damage to or destruction of the equipment or other property.

WARNING – Call attention to a potential danger that requires correct procedures or practices in order to prevent personal injury.



WARNING– To reduce the risk of electrocution:

1. Patients are not allowed to operate the product. Always unplug this product immediately while it's not in use.
2. Do not disassemble the Pump to avoid electrocution.
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 liquid. Do not use while bathing.
5. Do not reach for a product that has fallen into water. Unplug immediately.










WARNING–To reduce the risk of burns, electrocution, fire or injury to person:

1. The system must be operated with the mattress connected to the Pump. Please do not power-off or unplug the Pump while in use.
2. Always use the same voltage as stated on the label. Do not use other power cords on the Pump. Avoid children and pets acquire the plastic packing to prevent suffocation hazards.
3. Equipment is not suitable to use in the presence of a flammable anesthetic mixture with air or with oxygen or nitrous oxide.
4. Keep away from sharp objects.
5. Close supervision is necessary when this product is used by, on, or near children or people with disabilities.
6. Use this product only for its intended use as described in this manual. Do not use attachments not recommended by the manufacturer.
7. Never operate this product if the Pump has a damaged power cord or plug, if the Pump is not working properly, if the Pump has been dropped or damaged, or the Pump has been dropped into water. Return the product to a service center or to the distributor for examination and repair.
8. Keep the power cord away from heated surfaces.
9. Never block the air openings of this product or place the product on a soft surface, such as a bed or couch, where the openings may be blocked. Keep the air opening free of lint, hair, and other similar particles.
10. Never drop or insert any object into any air opening or hose tube.
11. Avoid dropping or putting any heavy object on the Pump.

12. Place the power cord and hose tube at patient's foot area to avoid strangling of patient's neck.
13. Avoid electromagnetic interference by not have strong electro-magnetic or RF generated equipment nearby patient's environment.
14. The Pump will have minor heat generated in operation, please do not direct contact the surface continuously for more than 1 minute.
15. When the mains supply is lost or had temporarily failed, the Pump will stop and the power failure alarm will alarm up to 20 minutes. This is normal and the product will return to normal operation once the mains supply is resumed.
16. The connection of the Lift-Up Detection to the Pump shall be performed by the operator. It is not designed for the patient to do the connection.
17. Connect this product to a properly grounded outlet only. See Grounding Instruction.
18. Avoid the risk of electrocution by grounding the ground pin with grounding power outlet.
19. The EMC specification is compliant with the regulation requirement. For power cord with ground pin (3 pin type), the connection with properly grounded power outlet would get a better EMC suppressing effect. The system will also work correctly for power cord connection without grounding.

Product Symbol Description

| SYMBOLS | DESCRIPTION |
|---|--|
| I | POWER ON |
| O | POWER OFF |
|  | ATTENTION |
|  | “BF” SYMBOL, INDICATE THIS PRODUCT IS ACCORDING TO THE DEGREE OF PROTECTION AGAINST ELECTRIC SHOCK FOR TYPE BF EQUIPMENT, APPLIED PART:MATTRESS |
|  | CAUTION, READ THE INSTRUTION BEFORE USE |
|  | AWAY FROM THE FLAME |
| IP21 | WATER AND DUST PROTECTION CLASSIFICAITON |
|  | FUSE SPECIFICATION |
|  | DISPOSAL OF ELECTRICAL & ELECTRONIC EQUIPMENT (WEEE): THIS PRODUCT SHOULD BE HANDED OVER TO AN APPLICABLE COLLECTION POINT FOR THE RECYCLING OF ELECTRICAL AND ELECTRONIC EQUIPMENT. |
|  | UL CERTIFICATION LOGO (COMPLIACE WITH IEC60601-1) WITH RESPECT TO ELECTRICAL SHOCK, FIRE AND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH IEC60601-1. |
| CB | CB CERTIFICATION LOGO |
| CE | CE CERTIFICATION LOGO |

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1. INTRODUCTION

This manual provides the information required for the initial set up and for the normal operation of the **Protekt® Aire 7000 Air Mattress System**. Before operating the **Protekt® Aire 7000 Air Mattress System**, be sure the operator had read and understood in detail the content of this manual.

2. INTENDED USE

The **Protekt® Aire 7000 Air Mattress System** is intended to help and reduce the incidence of pressure ulcers and provide the function of lateral rotation turning while optimizing patient's comfort. The **Protekt® Aire 7000 Air Mattress System** may be used in a variety of settings including, but not limited to individual home care setting and long-term care of whom suffering from pressure ulcer, or pain management as prescribed by physician.

The connection of Lift-Up Detection to the Pump shall be performed by the operator. It's not designed for the patient to perform the connection.



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

3. PRODUCT DESCRIPTION

The **Protekt® Aire 7000 Air Mattress System** is an alternating mattress replacement system used in the prevention and relief for patients with, or vulnerable to, pressure ulcers. The **Protekt® Aire 7000 Air Mattress System** offers patients a comfortable and relaxing support surface by using the established principles of alternating therapy, which can both prevent skin breakdown and enhance healing.

The Control Unit of the **Protekt® Aire 7000 Air Mattress System** is a Pump featuring a digital pressure adjustment function, mode selections, and audiovisual alarms. The 18 air cells mattress unit provides a unique design which keeps the lower sections of air cells constantly inflated while alternating and deflating the upper sections. The 3 static head cells remain static and provide a "pillow" support for optimum comfort. The mattress has a heavy-duty polyester base sheet with a quilted vapor permeable nylon with PU coated cover sheet.

Lateral rotation turning function is achieved by using special designed lateral rotation turning air cells to achieve the nursing needs. The lateral rotation turning function can function with alternate, pulsate, or static mode at the same time. The orientation of turning is defined by patient's orientation. Single turn mode can either maintain the

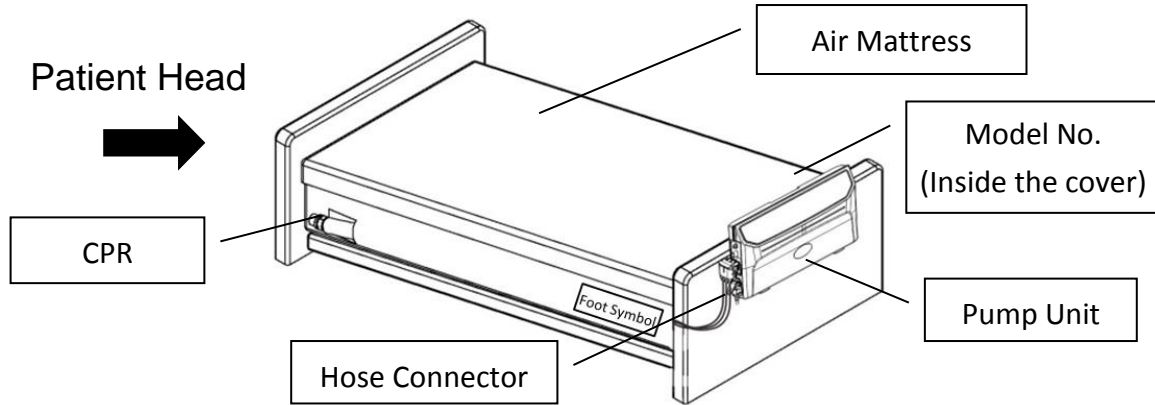
turning level constantly or alternate between turning and non-turning level. Full turn mode will alternate between left turn, horizontal level, right turn, and horizontal level periodically.

The system includes a rapid release twist CPR valve by the head section of the mattress for the event of cardiac arrest.

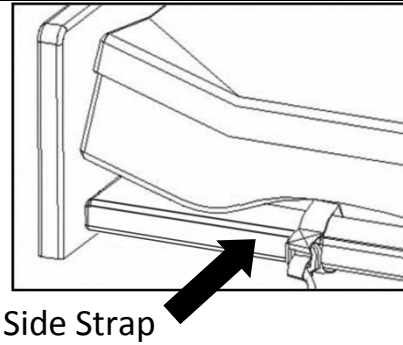
| Protekt® Aire 7000 Functions | | | | |
|------------------------------|---------|--------|--------------------|----------------|
| Main Function (Alternating) | | | Secondary Function | Other Function |
| Alternating | Pulsate | Static | Lateral Rotation | Fowler Mode |
| ■ | ■ | ■ | ■ | ■ |

4. PRODUCT INSTALLATION GUIDE

1. Unpack the box to inspect all items for any damage that may have occurred during shipping. If there is any damage, please contact your dealer immediately for assistance.
2. Place the mattress on top of the bed frame. The feet symbol on both side of the mattress indicates location of the foot end.

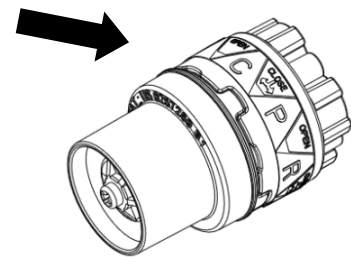


3. Secure the mattress onto the bed frame by using the elastic straps or binding side straps.

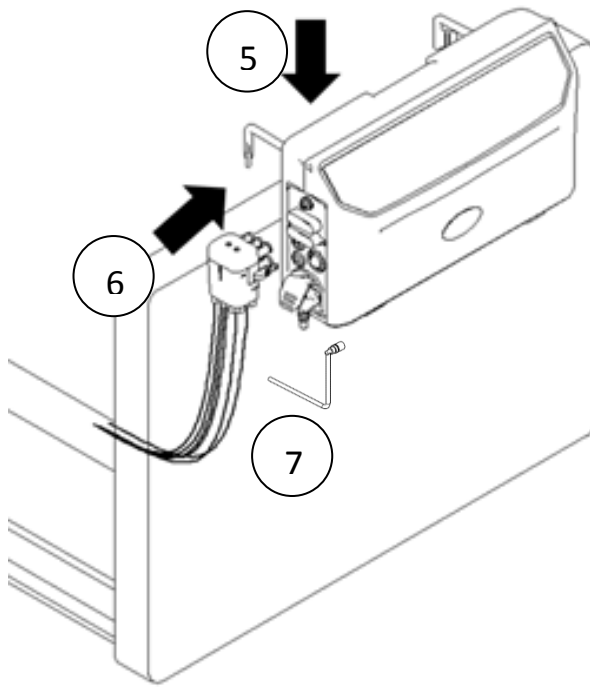


4. Ensure the CPR valve is at CLOSE position before turning on the power.

CPR Position Indicator

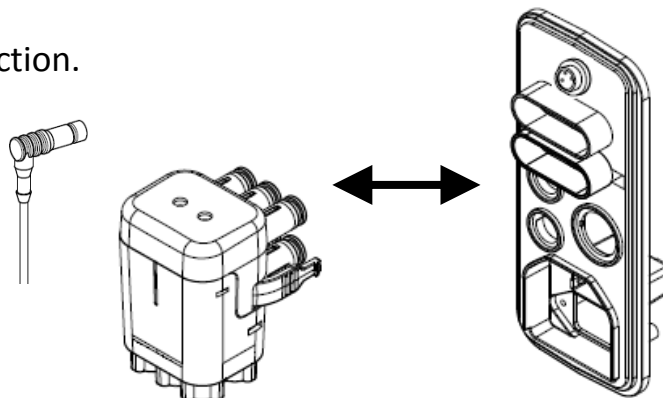


5. Position the Pump by its elastic hanger brackets over footboard of the bed. The elastic hanger brackets will self-adjust onto the footboard tightly.
6. Remove the Transport Cap of the hose connector and connect the hose connector to the Pump unit. Firmly push the hose connector into position and a “click” sound will secure the connection.
7. Connect the Lift-Up Detection cable to the Pump unit by first align and push-in the 3 pins connector then gently turn the screw clockwise to secure the connection.



WD433

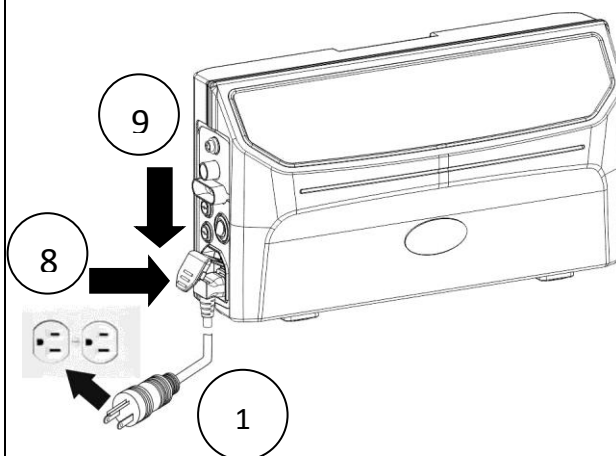
Follow the direction for connection.

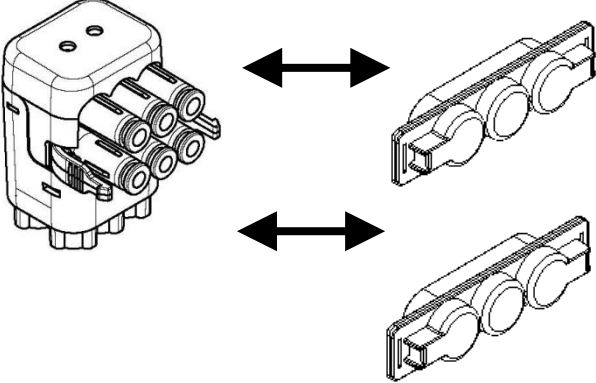


8. Connect the power cord to the Pump. The power switch should remain off.
9. Press the red power cord protector downward to secure the power cord.
10. Plug the power cord into the electrical outlet.

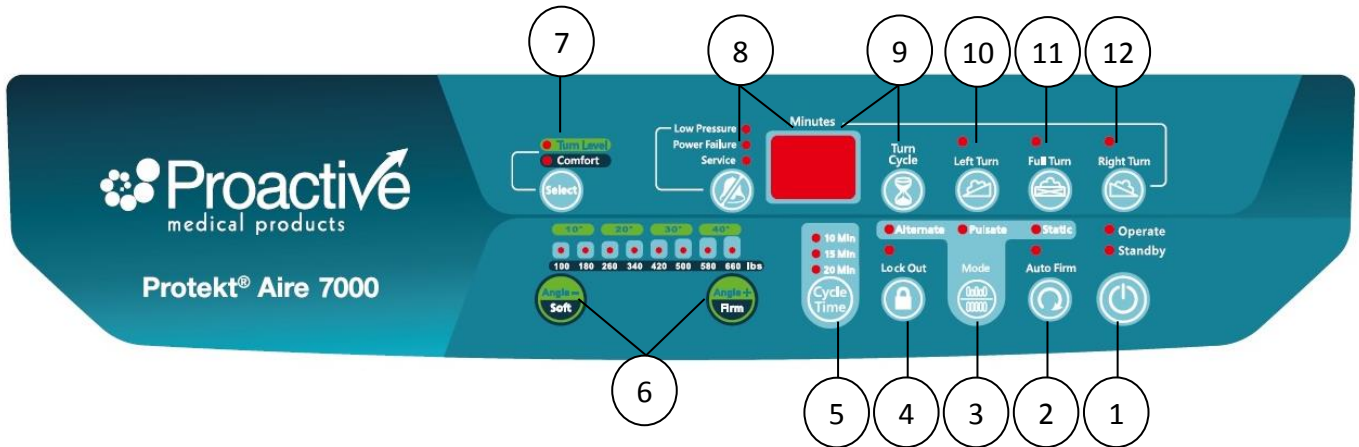
⚠ NOTE: Check and ensure the Pump unit is suitable for the local power voltage.

⚠ CAUTION: The Pump can only be applied to the mattress recommended by the manufacturer. Do not use the



| | |
|---|--|
| <p>Pump for any other purpose.</p> <p>⚠ WARNING: Do not place the Pump unit in area where power cord can come off easily.</p> | |
| <p>11. For patient's transportation, press "Auto Firm" button and wait for 5 minutes for the mattress to be inflated. Disconnect the hose from the Pump unit and put on the hose connector Transport Cap to keep the mattress inflated.</p> |  <p>Bi-directional Transport Cap</p> |

5. PANEL DISPLAY AND OPERATION GUIDE



5.1 PANEL DISPLAY

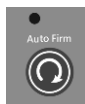
- ① Operate or Standby
- ② Auto Firm
- ③ Function Mode Selection (Alternate/Pulsate/Static)
- ④ Panel Lock-out
- ⑤ Alternate/Pulsate Cycle Time
- ⑥ Comfort Control (Soft and Firm)/Turn Level (Angle- and Angle+) Adjustment
- ⑦ Comfort and Turn Level select
- ⑧ Alarm Mute and Alarm Indicator
 - Low Pressure Alarm Indicator
 - Power Failure Alarm Indicator
 - Service (Malfunction) Alarm Indicator
- ⑨ Turning Cycle Time (Minutes) or Warning Code Display
- ⑩ Left Turn
- ⑪ Full Turn
- ⑫ Right Turn

5.1.1 Operate or Standby

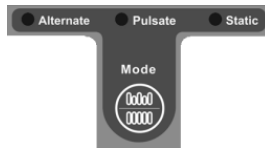


Press this button to start operating or go into standby.

5.1.2 Auto-Firm



The Pump will go into the inflation mode (LED lights flashing) every time the operate mode is triggered. This insures the mattress to be able to reach its maximum operating pressure. Once the max pressure level is reached, the Pump will automatically switch into the previous selected mode and comfort level. User can also use this function as full mattress inflation during patient sit-up or ingress/egress for better support.



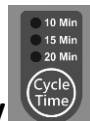
5.1.3 Function Mode Switch

- Alternate – the air cells of the mattress will be proportionally deflated to reduce the surface pressure. The alternating cycle will continue at the selected cycle time until another mode is selected.
- Pulsate –the mattress maintains at the selected pressure in static mode and changes from 100% to a reduced selected pressure according to the selected Pulsate cycle time. The Pulsate cycle will continue until another mode is selected.
- Static – the mattress maintains at a reduced selected pressure in alternate mode.



5.1.4 Panel Lock-Out

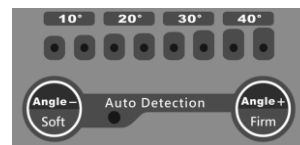
Press the Lock-Out button to lock the panel. Should the panel remain untouched for 30 seconds, the Lock-Out feature will lock the panel to prevent accident from changing setting without notice. To unlock, press the Lock-Out button for 3 seconds (LED light will flash while pressing the button).



5.1.5 Alternate/Pulsate Cycle Time Display

Alternate cycle time can be selected to 10, 15, or 20 minutes by pressing the Cycle button.

Pulsate cycle time can be selected to 10, 15, or 20 minutes by pressing the Cycle button.



5.1.6 Comfort Level and Turn Level Adjustment

Comfort level controls the air pressure output level. Press Firm button and the output pressure will increase and higher pressure output will support heavier weight patient, for decreasing air pressure, vice versa. Check to see if the suitable pressure is selected by sliding one hand between a deflated air cell and the patient’s buttocks areas and there should be minimum contact. Always leave at least 1 inch space between a deflated air cell and patient’s buttocks areas to prevent “bottoming-out”. Refer to **Table 1 Weight and Comfort Level Reference** for weight and comfort level suggestion.

Adjust the turning angle by select the Turn Level (refer to **5.1.7 Turn Level or Comfort Select**) and press Angle + to increase the turning degree angle or press Angle – to decrease the turning degree angle.



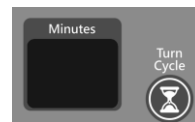
5.1.7 Turn Level or Comfort Select

Select turn level or comfort level by pressing this button.



5.1.8 Alarm Mute

Press the alarm mute button to suspend temporarily alarms. Should the situation not resolved and fault conditions continue, the alarm will resume notifying the patient and caregiver.



5.1.9 Turning Cycle Time (Minutes) Adjustment

- Single Turn (left or right turn): Press Turn Cycle to active periodic single turn.
- Full Turn (left and right turn alternatively): always periodic.
- Select turning time from 10-95 minutes by pressing the Turn Cycle button (the display time represents the continuing time for each turning position.)
- Hold the button for 3 seconds to reset the shortest turning time, 10 minutes.



5.1.10 Left Turn

- Initiate or cancel the Left Turn function by pressing this button.



5.1.11 Full Turn

- Initiate or cancel the Full Turn function by pressing this button.



5.1.12 Right Turn

- Initiate or cancel the Right Turn function by pressing this button.

5.2 OPERATION GUIDE


5.2.1 General Operation:





NOTE:






The power switch is located on the side of the Pump.


- Press  to turn on the unit, all LED indicators on the control panel will light up accompanied with a beep for 2 seconds (check for indicator failure if any), and the indicator of Standby on the control panel will light up. If the Pump was previously shut off in operate mode then the Pump will enter operate mode directly.



P.S. To test the battery, press  to turn off the power and the power failure alarm should be triggered. Refer to **5.2.3 Audiovisual Alarm** if the alarm is not triggered.




- Press the Operate button  and the system will begin to inflate and the "Auto-Firm" indicator will be flashing.
- The mattress should be fully inflated within 60 minutes, and automatically enter the previous operating mode, otherwise the low pressure alarm with warning code "L.F" will be triggered.

 Note: Do not proceed to other settings before inflation is completed.

- After initial inflation is completed. Press Auto-Firm button  for moving the patient onto the mattress. The mattress will turn into a steady condition in around 5 minutes. Move the patient onto the mattress and press Auto-Firm button  again to cancel Auto-Firm mode and select the appropriate mode.
- According to the weight of the patient, adjust the pressure setting to the most suitable level without "bottoming-out". User can determine an appropriate pressure by adjusting the Comfort Level. Please consult with your physician for a proper setting.

 Warning: the Pump unit should always be operating to prevent bedsores occurring.

- In operate mode, press operate/standby button  for the system to enter standby mode. The system should be in standby mode before shut down. Switch the power switch  to off and the warning code "S.d" will appear on the display to shut off the system.

 Note: For reminding purpose, power failure alarm will be triggered if the power is switched off in operating mode (refer to **5.2.3 Audiovisual Alarm**). Press power switch  to restart the system, or press Alarm Mute  to turn off the system (refer **5.2.4 Alarm Mute**).

- Activate or deactivate the lateral rotation turning by pressing the Turn button (refer to **5.1.10-5.1.12 Left, Full, and Right Turn**) after the panel has been

unlocked. If periodic lateral rotation is needed then press Turn Cycle button (refer to **5.1.9 Turning Cycle Time (Minutes) Adjustment**) for turn cycle adjustment.

Table 1 Weight and Comfort Level Reference

| Comfort Control (Indicators) | Patient Weight | | | | | | | | | |
|---------------------------------|----------------|-----|-----|-----|-----|-----|-----|-----|-----|------|
| | 132 | 165 | 209 | 264 | 330 | 418 | 528 | 660 | 800 | (lb) |
| | 60 | 75 | 95 | 120 | 150 | 190 | 240 | 300 | 380 | (kg) |
| | | | | | | | | | | |
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5.2.2 CPR

- When CPR needs to be performed, quickly rotate the CPR valve to “OPEN” position, at the same time, disconnect the hose connector from the Pump to speed up the air release.

5.2.3 Audiovisual Alarm

- Power Failure – When electrical shortage occurred or power cord is unplug without turning off the Pump or is pressed (intentionally or unintentionally), the “Power Failure” indicator will light up along with buzzer and will last 20 minutes.

NOTE: When the Pump has not been used for more than 3 months, the Pump may need 6 hours or more of operating time for the alarms to function properly.

- Low Pressure –When an abnormal low pressure occurred in body section for 2.5 minutes after the pressure is below the abnormal pressure threshold, the "Low Pressure" indicator will flash and beep. Should the situation not resolved and fault conditions continue, the alarm will resume.
- Service (Malfunction) –When fault conditions occur, the "Service" indicator will light up along with buzzer.

*Note: Refer to **Table 2 for Warning Code Reference Table** if error code appears on the display or refer to **10. TROUBLESHOOTING**.*

5.2.4 Alarm Mute

- When alarms are triggered, both LED light and buzzer will turn on to warn the patient and caregiver. By pressing the button, it will temporarily mute the buzzer so the caregiver may check for possible causes. Should the situation not resolved and fault conditions continue, the alarm will resume. Refer to **10. TROUBLESHOOTING** for diagnosis.
- During “power failure”, pressing “alarm mute” will cease all buzzers and indicators and turn off the system.
- During “low pressure alarm” if the pressure resumes back to normal then the low pressure alarm will stop.
- When more than one alarm is triggered, the alarm will be performed according to priority level. Refer to **Table 2 Warning Code Reference Table** for priority level.

Table 2 Warning Code Reference

| PRIORITY HIHG ↓ LOW | WARNING CODE | INDICATOR LED | AUDIBLE OUTPUT MODE | CONDITION OF OUTPUT | WARNING DESCRIPTION | REMARKS |
|------------------------------|-----------------|------------------|-------------------------------|--------------------------------|--|--------------------------------------|
| 0 | N/A | N/A | ONCE | Not in System Shutdown | Key Tone | Key Tone from Functional Button |
| 1 | | Power Failure | ONCE | POWER-OFF | System Shutdown | Shutdown |
| 2 | | ALL LED | ONCE | OPERATE OR STANDBY | Power-On | All Indicators Light On |
| 3 | N/A | N/A | ONCE | OPERATE OR STANDBY | State/Mode Switching | No Display |
| 4 | | Auto Firm | ONCE | OPERATE | Mattress Inflation Completion | Inflation Ended |
| 5 | | Auto Firm | ONCE | OPERATE | Auto-Firm Completion | Auto-Firm Ended |
| 6 | | Static | ONCE | OPERATE | Static Completion | Static Ended |
| 7 | N/A | Power Failure | REPEAT (Cycle 4 sec.) | POWER-OFF | Power Failure Alarm | No Display |
| 8 | | Low Pressure | REPEAT (Cycle 4 sec.) | OPERATE OR STANDBY | Power-On Inflation Failure Alarm | Inflate Failure |
| 9 | | Low Pressure | REPEAT (Cycle 4 sec.) | OPERATE OR STANDBY | Auto-Firm Failure Alarm | Auto-Firm Failure |
| 10 | | Low Pressure | REPEAT (Cycle 4 sec.) | OPERATE OR STANDBY | Low Pressure Overtime Alarm | Low Pressure |
| 11 | | Service | REPEAT (Cycle 4.5 sec.) | OPERATE OR STANDBY | High Pressure Overtime Alarm | High Pressure |
| 12 | | Service | REPEAT (Cycle 4.5 sec.) | OPERATE OR STANDBY | High Ambient Temperature Alarm | High Temperature |
| 13 | | Service | REPEAT (Cycle 4.5 sec.) | OPERATE OR STANDBY | Air Valve 1 Positioning Failure Alarm | Air Valve 1 failure |
| 14 | | Service | REPEAT (Cycle 4.5 sec.) | OPERATE OR STANDBY | Air Valve 2 Positioning Failure Alarm | Air Valve 2 failure |
| 15 | | Service | REPEAT (Cycle 15 sec.) | OPERATE OR STANDBY | Battery Low Alarm | Battery would need to be replaced |
| 18 | | NONE | NONE | OPERATE | LIFT-UP ANNGLE >45° (±15°) | Lift-up High |
| 19 | | NONE | NONE | OPERATE | LIFT-UP ANNGLE >20° (±15°) | Lift-up Low |
| 16 | | NONE | NONE | FACTORY CALIBRATION MODE | Calibration Not Completed | Calibration Unfinished |
| 17 | | NONE | NONE | FACTORY CALIBRATION MODE | Calibration Completed | Calibration Completed |

5.2.5 Lift-Up Detection

- “Auto-Firm” mode will be triggered when the upper half of the mattress is elevated and exceeded an angle of 45° ($\pm 15^{\circ}$) or more, and the warning code **LH** will appear on the display. The Pump unit will resume to previous setting when it’s lowered than an angle of 45° ($\pm 15^{\circ}$).
- Turning mode will be stopped if the upper half of the mattress is elevated and exceeded an angle of 20° ($\pm 15^{\circ}$) or more, and the warning code **LL** will appear on the display. If the angle is lower than 20° ($\pm 15^{\circ}$), the turning mode will be resumed.

6. CLEANING

Wipe the Pump unit with a damp cloth pre-soaked with a mild detergent, and keep the Pump unit away from dust. If other detergent is used, choose one that will have no chemical effects on the surface of the plastics case of the Pump unit.



CAUTION: Do not immerse or soak the Pump unit.

Clean the mattress cover by using single use wipes with a solution of neutral detergent and hand hot water. Rinse thoroughly with clean water and damp dry the mattress using single use wipes.

Disinfecting the cover

If the cover is heavily soiled or has been exposed to bodily fluids such as blood, it will require a more thorough cleaning procedure.

Use single use wipes with a 0.1% chlorine solution (1,000ppm) and cold water to wipe the cover. Rinse thoroughly with clean water and damp dry the mattress using single use wipes. Ensure the cover is completely dried before refitting to the mattress.

Frequent or prolonged exposure to higher concentration disinfectant solutions may prematurely age the fabric cover of mattresses. Cover surfaces should be protected during use and rinsed and dried thoroughly after disinfectant.

Laundering

- Before laundering, mattress cover should be completely removed.
- Remove Lift-Up Detection Sensor located on the bottom base before laundering.
- Mattress covers can be laundered as following:
 - Prewash 60°C +15 minutes
 - Main wash 60°C +15 minutes
- This should be followed by a cold rinse and extraction.
- Lift-Up Detection Sensor is not washable. It can be wiped as described, and install back after the mattress is completely dry.

Drying

Mattress covers should be hung from a line or bar and drip dried in a clean indoor environment. Covers must be completely dried before refitting to the mattress.

Mattress covers can be tumble dried on a low heat setting for 90 minutes. Drying temperature must not exceed 40°C. Exceeding the temperature can cause significant damage to the mattress cover.



CAUTION: Do not use phenolic-based product for cleaning.



CAUTION: After cleaning, dry the mattress without direct exposure of sunlight.

7. STORAGE

- To quickly vacuum air out from mattress for storage, rotate the CPR valve to OPEN position and disconnect the hose connector to release the air.
- Lay the mattress out flat and upsides down.
- Roll from the head end towards the foot end.
- Packing strap can then be stretched around the rolled mattress to prevent unrolling.
- The power cord could be wrapped around the Pump bumper or disconnected for storage.

8. MAINTENANCE

General

- Check main power cord and plug if there are abrasions or excessive wears.
- Check mattress cover for signs of wear or damage. Ensure mattress cover and tubes are stubbed together correctly.
- Check the air hoses for any kink or break. For replacement, please contact your local dealers.

Fuse replacement

- Disconnect the plug from mains power when a blown fuse is suspected.
- Remove the cover of the fuse holder by means of a small screwdriver.
- Insert a new fuse of the correct rating in, and replace the cover of the fuse holder back. The fuse rating should comply with the requested specification.

Air Filter Replacement


After checking **10. TROUBLESHOOTING**, if the air filter needs to be replaced:

- Replace the air filter located at the back of the Pump.
- The filter is reusable and can be washed gently with a mild detergent and water. Dry the filter before use.
- Check and replace air filter regularly if environment is dirty.

9. THE DISPOSAL OF AIR MATTRESS

When the air mattress is broken or no longer be useable, the mattress and the Pump may be discarded for recycle.


10. TROUBLESHOOTING

| PROBLEM | SOLUTION |
|--|--|
| The mattress is not able to connect with the Pump | <ul style="list-style-type: none"> ● Check if the mattress model (model no. located inside the cover by the foot end) xxAAxxx matches with the Pump model xxBBB-xxx. The AAA should be the same as BBB. If not, please contact with the agent or distributor. ● Check if the Transport Cap is removed and make sure the connector is not broken. |
| The Pump is showing no indications of working | <ul style="list-style-type: none"> ● Check if the plug is connected to the mains supply. ● Check if the power switch is switched to ON position (press ). ● Check if there is any blown fuse. |
| Power failure alarm failed | <ul style="list-style-type: none"> ● If the Pump is in operation but failed to trigger the power failure alarm during power off. Charge the Pump for 6 hours or more of operating time and if the power failure still not works then please contact the dealer or agent for further investigation. |
| The low pressure light is constantly flashing and the alarm is sounded | <ul style="list-style-type: none"> ● Check if the CPR is at CLOSE position ● Check if the connection between air tubes to Pump unit is tightly secured. ● Check if all coupling connections along mattress are secured. ● If the mains supply is normal but there is no sound of the Pump, please remove the connector from the Pump to check if there's air comes out. If not, please turn off the machine and contact the dealer or agent for further investigation. ● If all of above steps have been checked. Press "Alarm Mute" for system to be verified again. |
| The Pump is on but the mattress is not alternated | <ul style="list-style-type: none"> ● Ensure the mattress inflation is completed. ● Check the Pump control panel the indicator of "Alternate" should be lighted on, if not, switch it to "Alternate." ● Check if "Service" alarm indicator is on with buzzer, if yes, contact the dealer or agent for further investigation. |
| Service (Malfunction) Alarm is on | <ul style="list-style-type: none"> ● Press "alarm mute" for system to be verified again. If the alarm is still on, please contact dealer or agent. |
| The Pump is operating noisily | <ul style="list-style-type: none"> ● Make sure the Pump is resting against a solid surface. ● If the noise is getting louder, contact the dealer or agent for further investigation. |
| Patient is bottoming out (without alarm triggered) | <ul style="list-style-type: none"> ● Pressure setting might be inadequate for the patient, adjust comfort level to FIRM (refer to Table 1 Weight and Comfort Level Reference Table) and wait for a few minutes for better comfort. ● Follow the procedures "The low pressure light is constantly flashing and the alarm is sounded" for inspection. |

If the above information does not solve the problem, please contact your local dealer or agent for further support.

11. TECHNICAL DATA

11.1 Product Specification

| Pump Unit | | Air Mattress | |
|--|---|-----------------|--|
| Model | 80071 | Model | 8" Air Mattress (iLAL) |
| Dimension (cm) | 42.7(L)x 12.8(W)x 23.3(H) | Dimension (cm) | 200(L)x 90(W)x 20(H) 200(L)x 107(W)x 20(H) 30(H) with bolsters 40(H) Maximum turn level 50(H) Maximum turn level with bolster |
| Weight (kg) | 5.5 | Weight (kg) | 12 |
| Alter Cycle Time (minutes) | 10/15/20 | No. of Air Cell | 18 cells |
| Pulsate Cycle Time (minutes) | 10/15/20 | | |
| Turn Cycle Time (minutes) | 10—95 | | |
| Auto Firm Time (minutes) | 20 | | |
| Pump output flow range (LPM) | 9 + 9 | Cell Material | Nylon-TPU |
| Pressure Range(mmHg) | 25—60 (±5) | | |
| Input Voltage | AC120V 60Hz | Cover Material | Quilted nylon-PU |
| Input Current | 0.43 A _{MAX} (@132V~) | Bottom Material | Polyester-PU |
| Fuse Rating | T1A 250VAC | | |
| Classification | Class II with functional Earth | Max. Weight(kg) | 300 |
| | Type BF  | | |
| Warranty | 1 years | Warranty | 1 year |
| Environment Condition | | | |
| Operation Environment | 5°C—40°C 15%RH ~ 93%RH(No Condensation) | | |
| Storage Environment | -25°C—70°C ≦ 93%RH(No Condensation) | | |
| Environment Pressure | 70 kPa—101.3 kPa | | |
| Environment Level | ≦ 3000m | | |
| Water and Dust Protection Classification | IP21 | | |

11.2 EMC INFORMATION (120V)

| Guidance and manufacturer's declaration-electromagnetic emissions | | |
|--|-------------------|---|
| <p>The system is intended for use in the electromagnetic environment specified below.</p> <p>The customer or the user of the system should assure that it is used in such an</p> | | |
| Emission test | Compliance | Electromagnetic environment-guidance |
| RF emissions CISPR 11 | Group 1 | The system uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment. |
| RF emissions CISPR 11 | Class B | The system is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes. |
| Harmonic emissions IEC 61000-3-2 | Class A | |
| Voltage fluctuations /flicker emissions IEC 61000-3-3 | Compliance | |

Guidance and manufacturer's declaration-electromagnetic immunity

The system is intended for use in the electromagnetic environment specified below.

The customer or the user of the system should assure that it is used in such an environment.



| Immunity test | IEC 60601 test level | Compliance level | Electromagnetic environment-guidance |
|---|---|---|--|
| Electrostatic discharge(ESD) IEC 61000-4-2 | ± 6 kV contact ± 8 kV air | ± 6 kV contact ± 8 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 | ± 2kV for power supply lines ± 1kV for input/output lines | ± 2kV for power supply lines Not applicable | Mains power quality should be that of a typical commercial or hospital environment. |
| Surge IEC 61000-4-5 | ± 1kV line(s) to line(s) ± 2kV line(s) to earth | ± 1kV differential mode Not applicable | 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 | <5% UT(>95% dip in UT) for 0,5 cycle 40% UT(60% dip in UT) for 5 cycles 70% UT(30% dip in UT) for 25 cycles <5% UT(>95% dip in UT) for 5 s | <5% UT(>95% dip in UT) for 0,5 cycle 40% UT(60% dip in UT) for 5 cycles 70% UT(30% dip in UT) for 25 cycles <5% UT(>95% dip in UT) for 5 s | Mains power quality should be that of a typical commercial or hospital environment. If the user of the system requires continued operation during power mains interruptions, it is recommended that the system be powered from an uninterruptible power supply or a battery. |
| Power frequency(50/60 Hz) magnetic field IEC 61000-4-8 | 3 A/m | 3 A/m | The system power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment. |

NOTE UT is the a.c. mains voltage prior to application of the test level.

Guidance and manufacturer's declaration-electromagnetic immunity

The system is intended for use in the electromagnetic environment specified below.

The customer or the user of the system should assure that is used in such and environment.

| Immunity test | IEC 60601 test level | Compliance level | Electromagnetic environment-guidance |
|---------------------------------------|-------------------------------------|------------------|--|
| <p>Conducted RF IEC 61000-4-6</p> | <p>3 Vrms 150 KHz to 80 MHz</p> | <p>3 Vrms</p> | <p>Portable and mobile RF communications equipment should be used no closer to any part of the system including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.</p> <p>Recommended separation distance: $d = 1,2 \sqrt{P}$ $d = 1,2 \sqrt{P}$ 80MHz to 800 MHz $d = 2,3 \sqrt{P}$ 800MHz to 2,5 GHz</p> <p>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).</p> <p>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</p> <p>Interference may occur in the vicinity of equipment marked with the following symbol:</p>  |
| <p>Radiated RF IEC 61000-4-3</p> | <p>3 V/m 80MHz to 2,5 GHz</p> | <p>3 V/m</p> | <p>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</p> <p>Interference may occur in the vicinity of equipment marked with the following symbol:</p>  |

NOTE1 At 80 MHz and 800 MHz, the higher frequency range applies.

NOTE2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

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

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

**Recommended separation distance between
portable and mobile RF communications equipment and the system**

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

| Rated maximum output power of transmitter W | Separation distance according to frequency of transmitter m | | |
|---|--|--|---|
| | 150 kHz to 80 MHz $d = 1,2\sqrt{P}$ | 80 MHz to 800 MHz $d = 1,2\sqrt{P}$ | 800 MHz to 2,5 GHz $d = 2,3\sqrt{P}$ |
| 0,01 | 0,12 | 0,12 | 0,23 |
| 0,1 | 0,38 | 0,38 | 0,73 |
| 1 | 1,2 | 1,2 | 2,3 |
| 10 | 3,8 | 3,8 | 7,3 |
| 100 | 12 | 12 | 23 |

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

NOTE1 At 80 MHz and 800 MHz, the separation distance for the higher frequency range applies.

NOTE2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

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