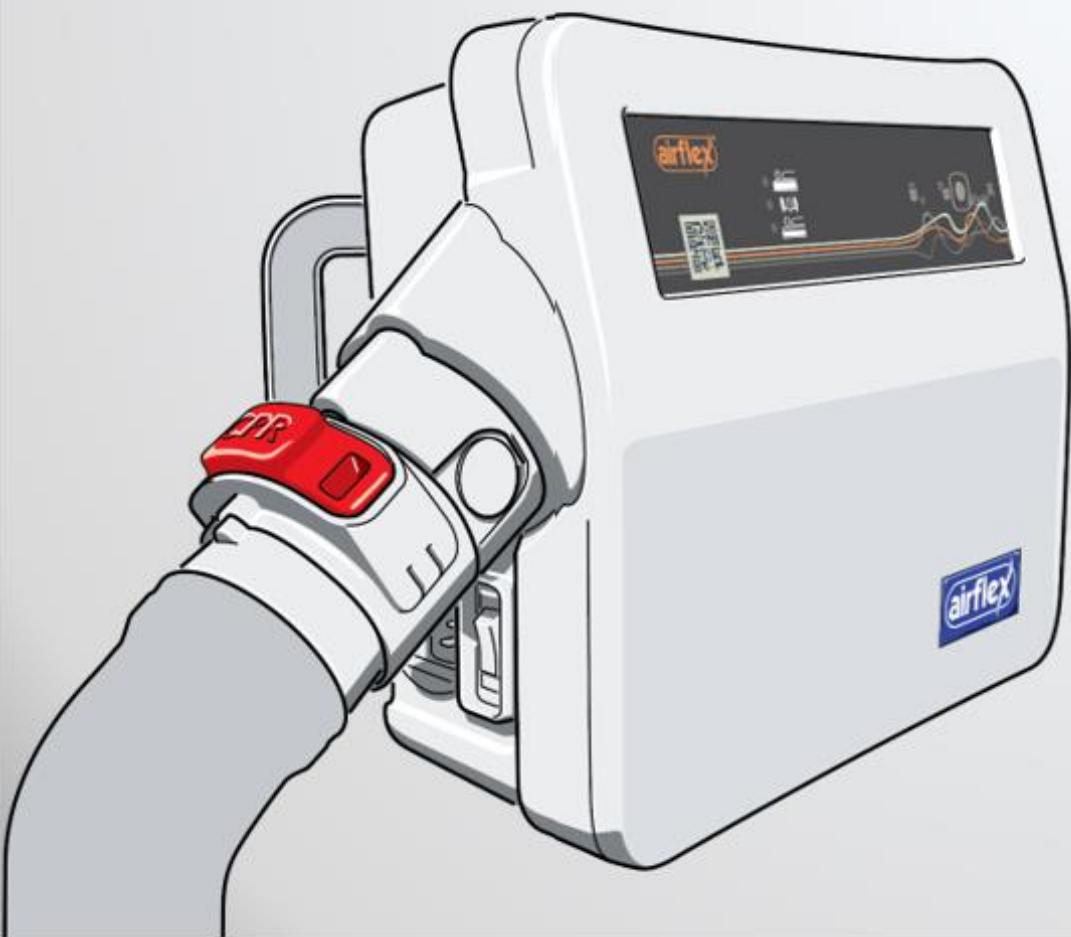


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airflex[®]
Overlay

**USER INSTRUCTION
MANUAL**





USER INSTRUCTION MANUAL

INSTRUCTIONS FOR USE

Version 4.0

Sep 2025



Version History

Version	Implemented By	Revision Date	Approved By	Approval Date	Reason
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1.1	Ben Smith	June '15	Graham Ball	June '15	Reformatting, Amendment to Hooks, pg 10
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Contents

Introduction.....	7
Incident Reporting.....	7
Intended Use	8
Further Information	9
Contraindications	10
Descriptions of Symbols	11
List of Components.....	12
General Cautions / Warnings	13
Installation.....	15
Display Screens / Operation Modes	19
Loading	19
Lockout Function	19
Automatic Alternating Mode	19
Different Modes / Displays.....	20
Firm Alternating	22
Firm Static.....	22
Soft Alternating	23
Soft Static	23
Alarms.....	24
High Pressure Alarm	24
Low Pressure Alarm.....	25
Power Failure Alarm	25
CPR Emergency.....	26
Transportation Mode	27
Preparation.....	27

Storage (and Shipping)27

Cleaning and Maintenance.....28

 Cleaning Procedures.....28

 Cleaning the mattress28

 Cleaning the Pump Unit.....30

Technical Parameters31

 Power Unit.....31

 Mattress31

 Internal Components Operating Conditions31

Introduction

Congratulations on purchasing the Airflex® OVERLAY three-cell cycle alternating pressure air mattress. This high quality medical product, if installed and cared for in accordance with these instructions, will provide many years of trouble free service and outstanding clinical performance.

The Airflex® OVERLAY is an alternating pressure mattress replacement system designed for the prevention of pressure ulcers.

This mattress system is manufactured to conform with international standards, EN 60601-1, EN 60601-1-2, and is registered to CE & UKCA.

Please read this Instruction Manual thoroughly before using the Airflex® OVERLAY.

Incident Reporting

In the event of a serious incident occurring in relation to this device, this should be reported to the manufacturer (Rober Ltd) and the competent authority of the member state in which you are situated.

Intended Use

The Airflex® OVERLAY is a two-cell cycle alternating pressure air mattress replacement system. It is designed to prevent and treat pressure sores for patients who are at low to medium risk of developing them. It also aids the healing of pressure sores in patients up to Grade 2 pressure injuries.

The objective of the Airflex® OVERLAY is to reduce the contact pressure created by the patient lying on the mattress. This is achieved by sequentially inflating and deflating the air cells of the mattress. This regularly reduces the contact pressure which restores blood flow and tissue oxygenation in the deep tissues about the vulnerable bony prominences.

The mattress system can fit onto any bed type in hospitals, nursing homes and in the community. This mattress replacement system is quiet, comfortable and simple enough for a single caregiver installation. All operator controls have been deliberately minimised to improve user interaction.

The system consists of an air filled mattress that is placed on top of a conventional bed frame and is connected via an air supply hose to a separate microprocessor controlled Power Unit.

The Power Unit is software driven using a solid-state pressure sensor so it has the capability to monitor the air pressure in the mattress continuously. The system is completely self regulating and automatically alters the air pressure depending on the weight of the patient and their position in bed.

An emergency deflation valve system is incorporated to allow cardiac resuscitation procedures. Named appropriately as a CPR

unit, it also has self sealing valves which allow the mattress to remain inflated for 24 hours when disconnected from the system.

A number of audio and visual alarms exist to warn caregivers of any problems within the system.

N.B. As with all alternating mattress systems, the Airflex® OVERLAY is contraindicated for use where patients have unstable fractures or spinal injuries.

Further Information

The Airflex® OVERLAY sits atop any hospital mattress. It is quiet, comfortable and simple enough for a single caregiver installation.

The frequent relief of contact (interface) pressure has been demonstrated to reduce the risk of pressure injury which is caused by reduced capillary blood flow in the tissue surrounding the bony prominences. The **Airflex® OVERLAY** is also a valuable aid in the treatment of existing sores (ulcers).

The objective of the Airflex® OVERLAY is to relieve the contact pressure created by the patient lying on the mattress overlay. This is achieved by sequentially inflating and deflating the air cells of the overlay. This regularly relieves the contact pressure which restores blood flow and tissue oxygenation in the deep tissues about the vulnerable bony prominences.

The system consists of an air overlay that is placed on top of a conventional mattress and is connected via an air supply hose (umbilical) to a separate control unit. This overlay mattress

consists of a single deck of 18 transverse air cells, protected by the cover.

The control unit is software driven which gives the capability to maintain the air pressure in the cells at the required level and to automatically alter the air pressure depending on the weight of the patient and their position in bed.

The additional presence of a CPR valve located at the side of the unit permits the rapid deflation of the mattress in an emergency.

Contraindications

As with all alternating mattress systems, the Airflex® OVERLAY is contraindicated for use where patients have unstable fractures or spinal injuries.

Descriptions of Symbols

Throughout this user manual and on the pump unit itself, the Airflex® OVERLAY has numerous symbols:

Symbol

Explanation



Type B – Applied Part



Class II Medical Device (certified to Class 1)

O / I

Off / On



Refer to Instruction Manual



General Warning



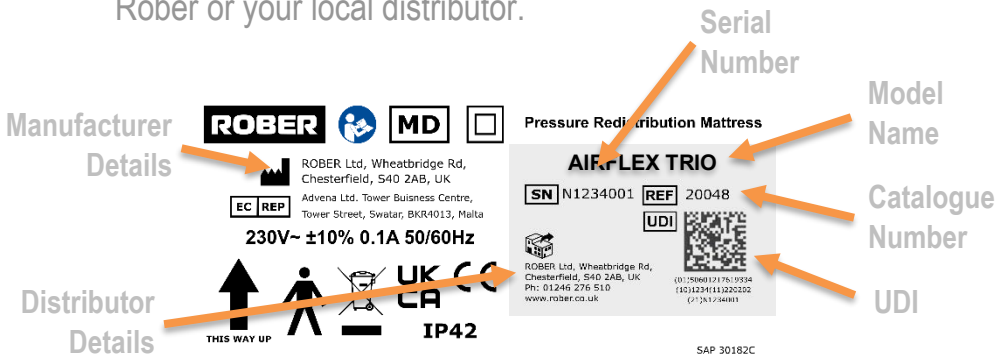
Mode

Button (for operation of pump unit only)

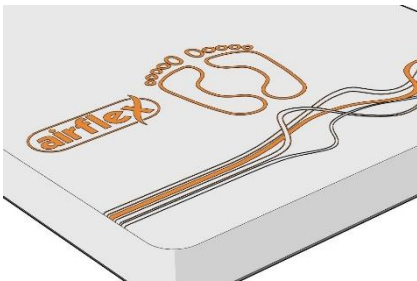
List of Components

Unpack your Airflex® OVERLAY mattress system and check that you have the following:

- Airflex® OVERLAY Pump Unit:
- Please note the Name, Model and the Serial Number of your pump in case you need to reference these when contacting Rober or your local distributor.



- Airflex® OVERLAY Mattress:



The top cover is made from Dartex™ which is an antimicrobial, multi-stretch, vapour permeable, PU coated fabric. The bottom cover is made from durable PVC coated Nylon.

- Power Cord:

The power plug should be appropriate to your country or region.

General Cautions / Warnings



Before setting up the mattress, it is important to read all of the safety instructions below. Any actions or operations performed outside the guidance of this manual are carried out at the user's risk. Rober and/or the supplier are not liable for any such actions.

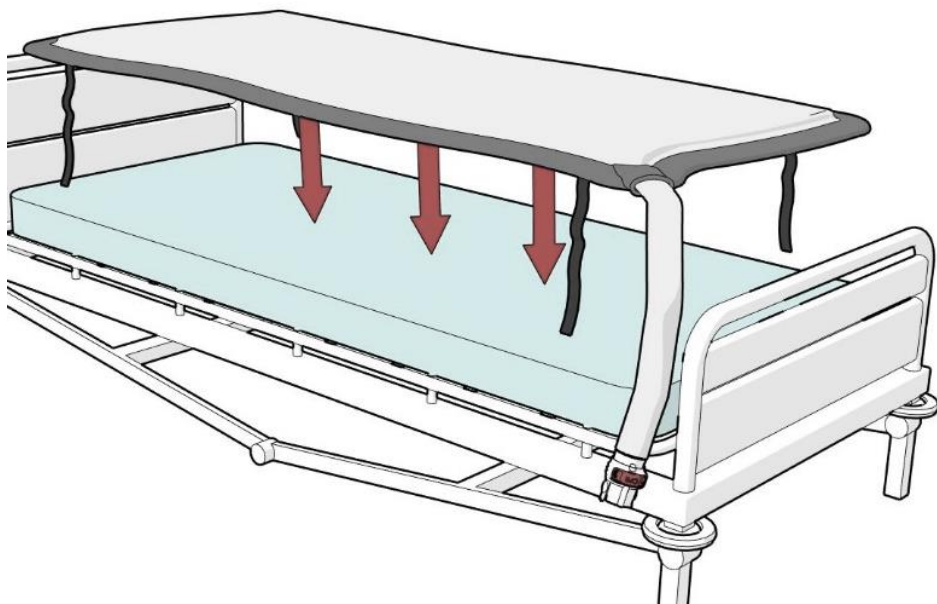
- Inspect before use. If any faults are discovered, the mattress system should not be used and the supplier immediately notified.,
- Check the power cord is positioned so there is no risk of obstruction or potential injury.
- **Risk of explosion:**
Do not use in the presence of flammable chemicals or gases.
- **Risk of fire:**
Do not place near heat sources or naked flames.
- **Risk of electrocution:**
Ensure that the mains power is fully earthed.
Never partially or fully submerge electrical equipment in fluids.
Check the power cord is free from any damage.
Never handle the plug with wet hands.
Do not open the case. Any repairs should be made by qualified service engineers.
Do not spill food or liquids on the pump unit. *If spillage occurs, disconnect from the mains and allow to dry before re-use.*
- Do not use phenol based cleaning solutions on the mattress or cover.

- Do not place any components of the mattress system in either heat or steam autoclaves.
- **Risk of Suffocation:**
The cover is not air permeable. Ensure the patient can use the product safely.
- Do not drop the pump unit and do not drop objects onto it.
- Never use sharp objects on or near the mattress.
- Never insert objects into the pump.

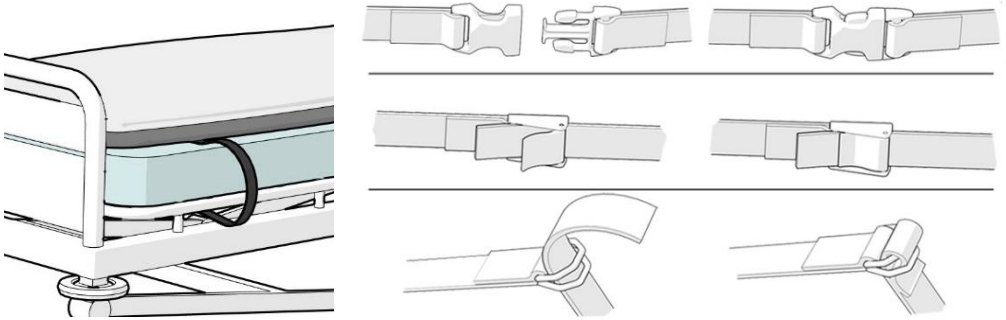
Installation

Please follow these instructions to install your Airflex® OVERLAY mattress overlay safely and appropriately.

1. Place the mattress overlay atop a conventional mattress, ensuring the light grey top cover is facing upwards and the air supply hose (umbilical) is positioned at the foot end of the bed.

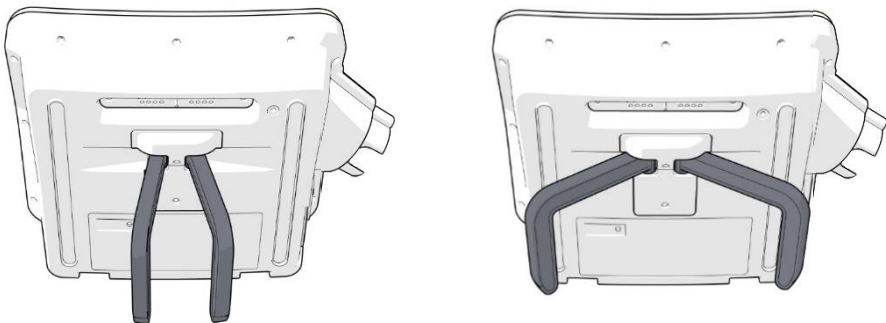


2. Secure the mattress to the bed using the straps and fasteners provided.

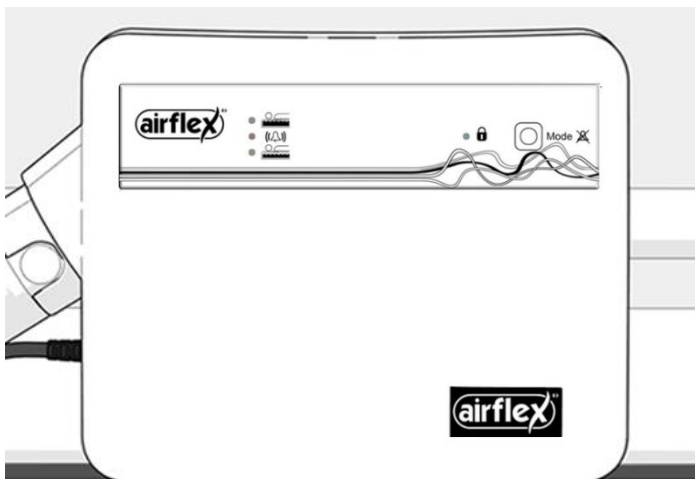


Ensure that fastened straps do not restrict the operation of any moving parts of the the hospital bed. Straps attached around both moving parts and fixed parts of a bed frame will cause severe damage to the mattress. After installation carefully test the functions of the bed to ensure the mattress moves freely with the bed plattform.

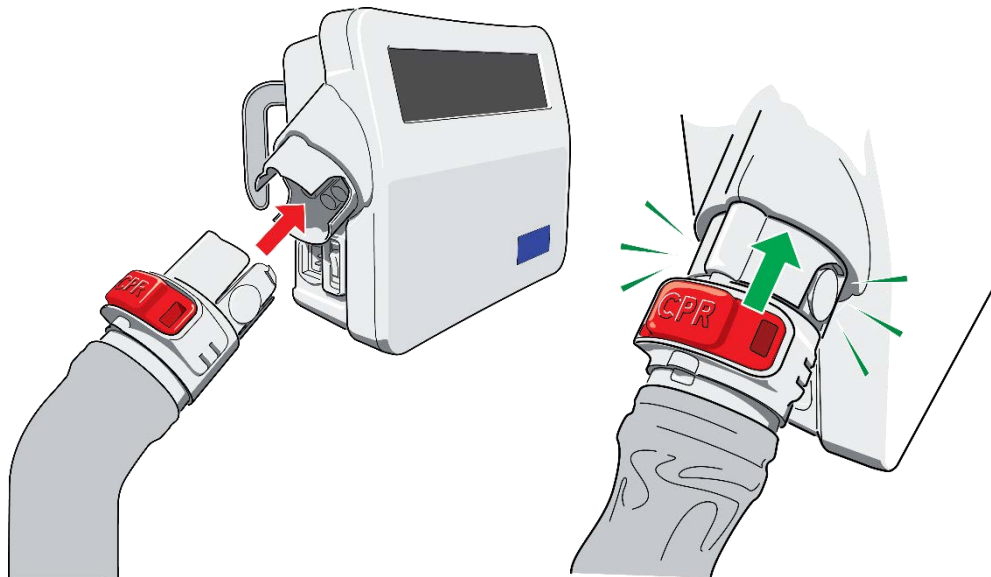
3. The hooks on the back of the Airflex® OVERLAY have been designed to accommodate a variety of bed frames and can be adjusted easily by opening out the hooks and rotating. Once released they will clamp to the bed frame using spring pressure.



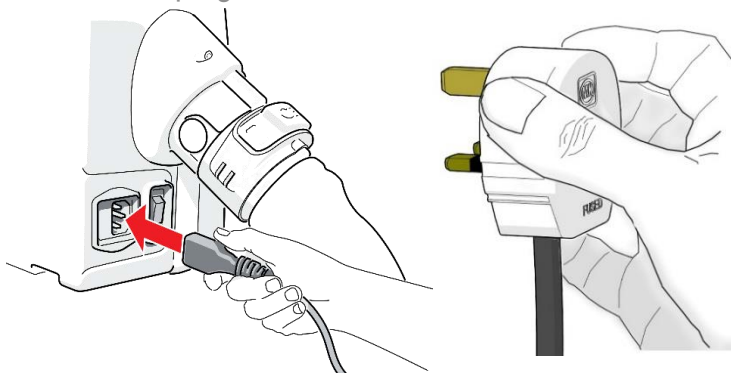
4. Hang the unit onto the bed frame at the foot end of the bed with the control panel facing outwards.



5. Check that the CPR clip is in the closed position and plug the umbilical hose into the dock.



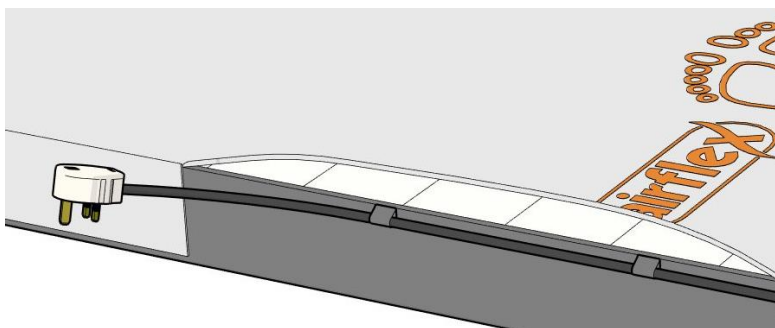
6. Plug the power cable into the power cable into the pump using the international connector and then plug the other end (*the main disconnection devise*) into an **easily accessible** mains supply socket. Turn the unit on using the switch to the right of the unit's international plug socket.



Due to the special design on the power cable's connection, it cannot be pulled out easily without pushing the yellow button. Ensure that the pump unit is positioned so this button is accessible.



Ensure that the power cable is safely guided through the loops located under the skirt (waterfall). This will help reduce any hazards in relation to the power cable.



7. Wait for approximately 15-20 minutes and your **Airflex® OVERLAY Mattress System** is ready for use.

Display Screens / Operation Modes

Before using your Airflex® OVERLAY it is important to understand the display on the front panel of the pump unit and the various operation modes of the system

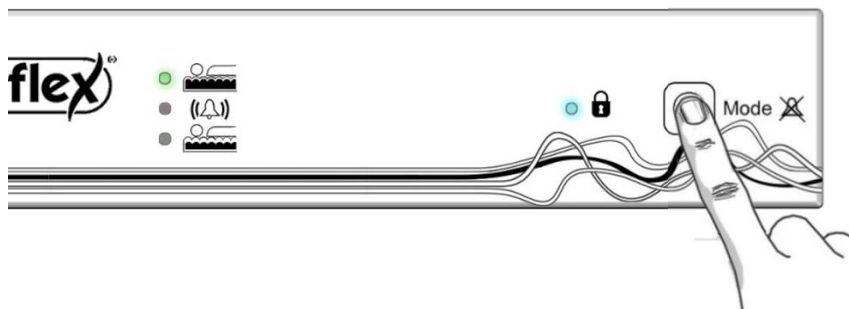
Loading

Whilst the Airflex® OVERLAY is inflating the mattress two GREEN LEDs, flash alternately.



Lockout Function

60 seconds after either full inflation or mode change, the system will automatically lock out the display panel.



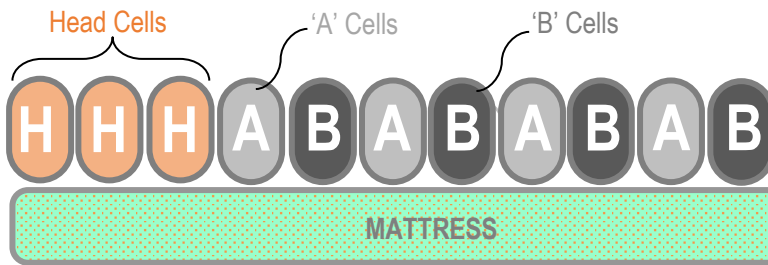
This is indicated by the blue LED next to the padlock symbol. To unlock the front panel hold down the "Mode" button until the Blue LED goes out.

Automatic Alternating Mode

Once the unit has inflated the mattress it will automatically default to Firm Alternating Mode (see below) and calibrate to the patients weight.

Different Modes / Displays

The 18 air cells of the Airflex® OVERLAY mattress are pneumatically connected into three groups.

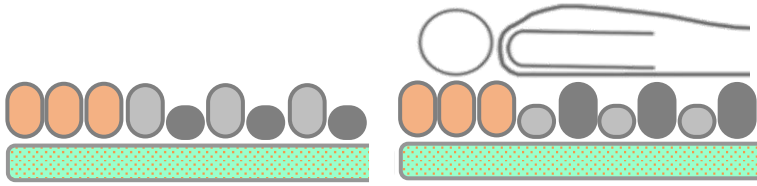


The 3 air cells at the head of the mattress ("head" cells) remain permanently inflated.

The remaining 15 air cells inflate and deflate in a 1 in 2 pattern with a cycle time of 10 minutes.

When the "A" cells deflate, pressure is relieved under that portion of the body in contact with those cells.

After 5 minutes, the "A" cells re-inflate and the "B" cells deflate for the next 5 minutes.



At the end of 10 minutes, contact pressure is cyclically offloaded from all parts of the body in contact with the active components of the mattress overlay.

Airflex® OVERLAY has four modes:

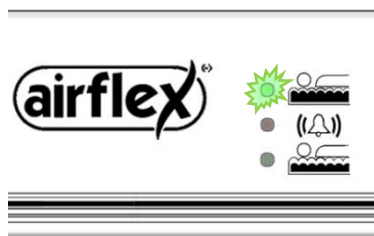
- Firm Alternating
- Firm Static
- Soft Alternating
- Soft Static

Press the mode button to cycle through these different modes, until the desired mode is found.

Firm Alternating

TOP GREEN LED – **FLASHING**

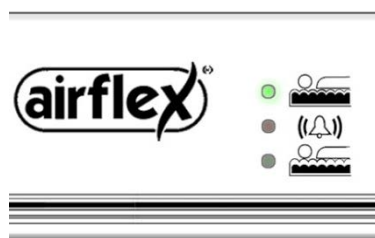
On start-up the system defaults to Firm alternating mode and automatically calibrates to patient weight and body position.



Firm Static

TOP GREEN LED – **STEADY**

The mattress calibrates to patient weight and remains in static mode for 30 minutes. In static mode all cells are inflated.



Soft Alternating

BOTTOM GREEN LED – **FLASHING**

The mattress changes to a lower range of pressure settings more suited to the comfort of small and under weight patients.



Soft Static

BOTTOM GREEN LED – **STEADY**

The mattress remains at the lower range of pressure settings and alternation ceases. The Airflex OVERLAY reverts back to 'Soft Alternating' after 30 minutes.



The 'SOFT' setting of the Airflex® OVERLAY is suitable for frail underweight patients and children.

N.B. *Both Static Modes revert to Alternating Mode after a preset period of 30 or 120 minutes, depending on your local distributor.*

Alarms

Each of the following alarm displays will be accompanied by a fixed tone audible alarm. The audible alarm can be cancelled by pressing the “Mode” Button, however if the cause of the alarm is not resolved after 15 minutes the audible alarm will reactivate.

To turn off the alarm permanently, or reset the pump after fixing the cause of the alarm, simply switch the pump unit off and then on again.



THE PUMP MUST BE RESET AFTER ANY ALARM EVENT.
ALL ALARM CONDITIONS ARE SHOWN WITH A RED LED

High Pressure Alarm

This condition could be caused by an over pressure in the mattress e.g. by another person sitting on the mattress. This alarm is shown by a **FAST FLASHING RED LED**.

To resolve this issue, unclip the CPR clip and let out a small amount of air and then reseal the CPR clip. Reset the pump afterwards. Persistent high pressure alarms may indicate an internal fault. If this occurs Care should cease. The pump should immediately be sent for servicing.

Low Pressure Alarm

This condition could be caused by releasing the CPR clip or a leak in the mattress. This alarm is shown by a **SLOW FLASHING RED LED**.

Power Failure Alarm

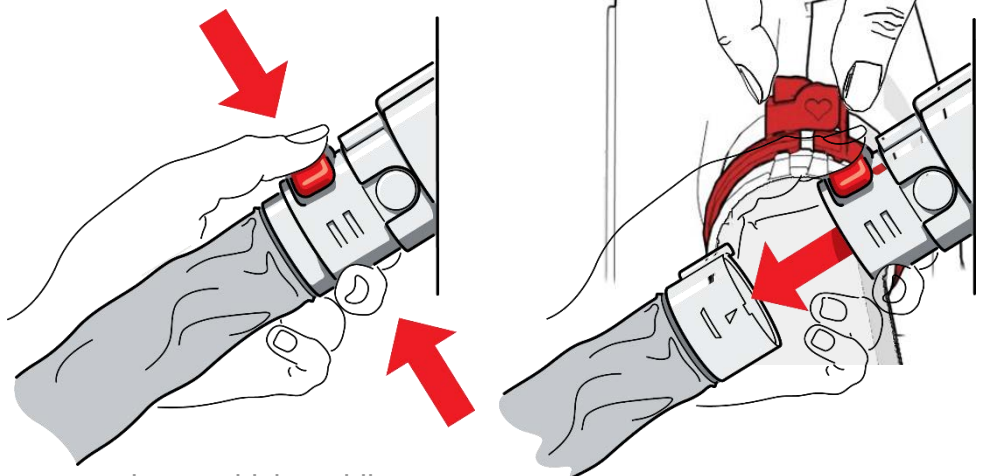
If there is a failure of the mains power supply or the unit is accidentally unplugged from the mains then a **CONTINUOUS RED LED** will show.

N.B. *The Power Failure Alarm is powered through a small battery.*

The sole function of this battery is to operate the alarm in case of a mains power failure.

CPR Emergency

In the case of a patient emergency where CPR may be necessary, the Airflex® OVERLAY has been equipped with a CPR



release which rapidly deflates the mattress so that cardiac resuscitation can be performed on a firm surface.

To deploy this emergency system, simply grip the CPR unit and push the RED BUTTON. The hose end closest to the mattress should spring out and the mattress will rapidly deflate.



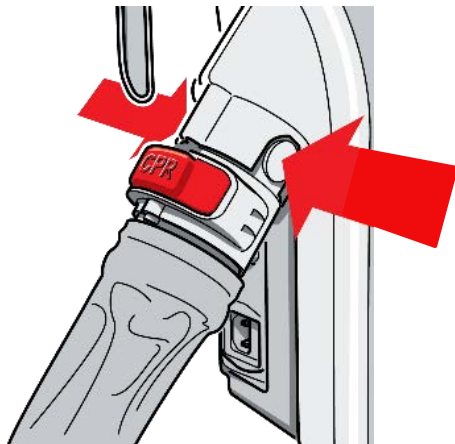
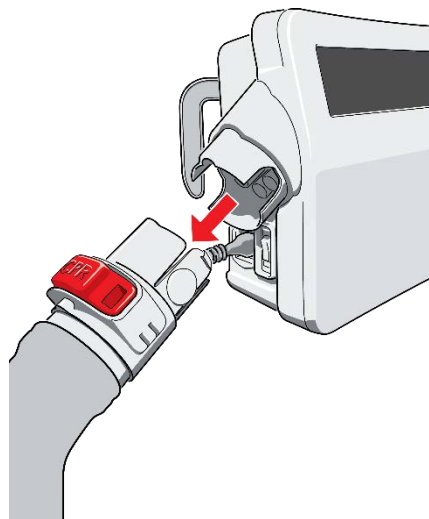
Ensure that after the CPR release has been disconnected, and CPR is complete, the hose end is pushed back in until an audible click is heard from the Red Button and then the pump is reset.

Transportation Mode

In this configuration the mattress can be transported easily around the hospital.

Preparation

When separating the pump unit from the mattress the unit should first be SET TO STATIC MODE.



When all cells are inflated the pump should be SWITCHED OFF and the umbilical hose separated

Simply push the buttons on the sides of the umbilical dock and the umbilical will disconnect. The mattress itself will remain inflated for several hours.

Storage (and Shipping)

The Power Unit should be stored standing upright in a vertical manner NOT laid flat. The following conditions should apply, where possible, for the longevity of internal components:

Ambient Temperature	+10°C to 40°C
Relative Humidity	10% – 100% non condensing
Atmospheric Pressure	700hPA – 1060hPA

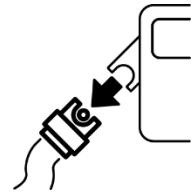
Cleaning and Maintenance

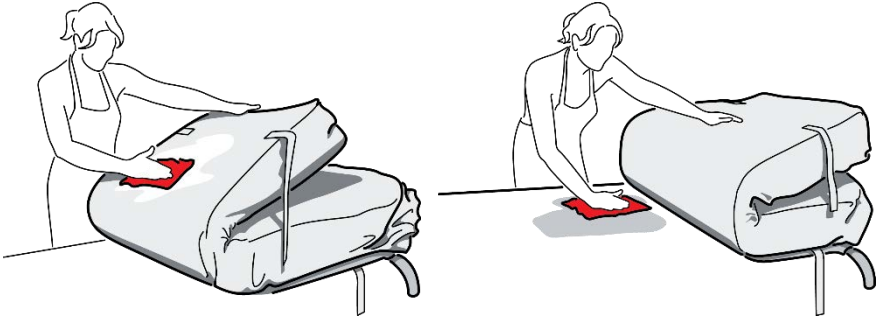
Cleaning Procedures

Cleaning the mattress system every week is recommended, or sooner if heavily soiled. Please make sure that hands are cleaned and appropriate protective clothing is worn before commencing the cleaning process.

Cleaning the mattress

- 1. Leave the cover on the mattress and disconnect the mattress from the unit.*
- 2. Clean the surface of the wash-down table with hypochlorite solution or an equivalent disinfectant.*
- 3. Wash mattress top cover using hot water (60 °C) and detergent. Dry with a paper towel.*
- 4. Fold mattress in half-length ways and clean both the table and the exposed mattress.*





5. Turn over mattress so the other bottom half is exposed and clean both mattress and table.



6. Repeat Steps 3 – 5 using hypochlorite solution or equivalent disinfectant.



7. Wipe the umbilical hose with hot water and detergent. Dry with a paper towel and repeat wiping down using hypochlorite or equivalent solution.



*If required, the cover can be removed and machine-washed at a temperature of **80 - 90 °C for a minimum of 10 minutes.***

The cover may also be sterilised using ETO or Draeger processes.

Cleaning the Pump Unit

The Pump Unit can be cleaned by wiping a cloth dampened by a disinfectant solution or sterilised using ETO or Draeger processes.



Ensure the unit is disconnected from the mains electricity supply before cleaning.

DO NOT autoclave the mattress or pump unit.

Maintenance



Only qualified technicians, trained or formally approved by Rober Limited, may carry out maintenance, modifications or repairs on the mattress system.



NO maintenance or servicing should take place while the Mattress System is in use. Ensure that the patient is clear of the system when carrying out maintenance.

Technical Parameters

Power Unit

Model	Airflex® OVERLAY
Dimensions (L x W x H)	400 x 160 x 250 mm
Weight	2.8 Kg
Power Supply	230V~ ±10%, 50 / 60 Hz
Fuses	2 x T1.6A, 250V~ 5 x 20mm
Power Consumption	20w max
Ingress Protection Rating	IP42
Classification	Class B Medical Device Class II (certified to Class 1)
Operating Cycle	10 minute (Continuous)
Flammability Rating	UL94 V0

Mattress

Model	Airflex® OVERLAY
Dimensions (L x W x H)	1900 x 880 x 120 mm (<i>Standard</i>) 1900 x 800 x 120 mm (<i>Narrow</i>)
Number of Air Cells	18
Weight	6.7 kg
Mattress Material	Polyurethane
Flammability Rating	Complies with BS7175 Ignition Source 0, 1 and 5
Maximum Weight of Patient	180kg

Internal Components Operating Conditions

Ambient Temperature	+10°C to 35°C
Relative Humidity	30% – 75% non condensing
Atmospheric Pressure	700hPA – 1060hPA

Your local distributor is:-



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