

HME SIGNATURE SERIES DOVE MATTRESS

USER MANUAL





DISCLAIMER:

The User Manual is based on general product applications. Specific clinical assessments should be based on individual client physical and functional assessments by a healthcare professional.

Manufactured By: Air Kinetic Technologies Corp.

Distributed By: HME Home Health Ltd. (www.HMEBC.com, Info@hmebc.com)

IMPORTANT SAFEGUARDS

READ ALL INSTRUCTIONS BEFORE OPERATING THIS DEVICE

NOTE - CAUTION AND WARNING STATEMENTS:



- NOTE Indicates a tip.
- **CAUTION** Indicates correct operating or maintenance procedures to prevent damage to or destruction of, the equipment or other property.
- WARNING Indicates a potential danger that requires correct procedures or practices to prevent personal injury.

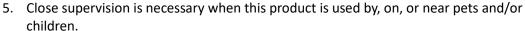
WARNING – To reduce the risk of electrocution:

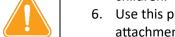


- 1. Always unplug this product immediately when not in use.
- 2. Do not disassemble the Control Unit.
- 3. Do not place or store the product where it can fall or be pulled into a tub or sink.
- 4. Do not place in or drop into water or other liquid. Do not use it while bathing.
- 5. Do not reach for a product that has fallen into the water. Unplug immediately.

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

- 1. The operation of this system requires that the mattress is connected to the Control Unit. Please do not power off or unplug the Control Unit while in operation.
- 2. Always use the same voltage as stated on the label. Do not use other power cords on the Control Unit.
- 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.





- 6. Use this product only for its intended use as described in this manual. Do not use attachments that are not recommended by the manufacturer.
- 7. Never operate this product if the Control Unit has a damaged power cord or plug, if the Control Unit is not working properly, if the Control Unit has been dropped or damaged, or if the Control Unit has come in contact with water. Return the product to a service center or distributor for assessment 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 Control Unit.



- 12. Place the power cord and hose tube at the foot of the bed to avoid tripping or other hazards with the cord.
- 13. Remove all electromagnetic or RF-generated equipment from close proximity, to avoid electromagnetic interference.
- 14. The Control Unit will have minor heat generated in operation, avoiding prolonged contact.
- 15. When the main power supply is lost or has failed temporarily, the Control Unit will stop working and the power failure alarm will sound for up to 20 minutes. This is normal and the product will return to normal operation once power is resumed.

PRODUCT SYMBOL DESCRIPTION

I	POWER ON
0	POWER OFF
\triangle	ATTENTION
	DOUBLE INSULATION
†	THE "BF" SYMBOL, INDICATES THIS PRODUCT IS IN ACCORDANCE WITH THE DEGREE OF PROTECTION AGAINST ELECTRIC SHOCK FOR TYPE BF EQUIPMENT, APPLIED PART: MATTRESS
	CAUTION, READ THE INSTRUCTION MANUAL BEFORE USE
	KEEP AWAY FROM FLAMMABLE MATERIALS
IP21	WATER AND DUST PROTECTION CLASSIFICATION
T5A 250V	FUSE SPECIFICATION
A	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.
C UL US	UL CERTIFICATION LOGO (COMPLIANCE WITH IEC60601-1) WITH RESPECT TO ELECTRICAL SHOCK, FIRE AND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH IEC60601-1.



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1. Introduction

This manual provides the information required for the initial setup and normal operation of the **HME Signature Series Dove Micro Low Air Loss Mattress System**. Before operating this Mattress System, be sure the operator has read and understood in detail the content of this manual.

2. Intended Use

- The **HME Signature Series Dove Micro Low Air Loss Mattress System** is intended to reduce the incidences of pressure wounds while optimizing an individual's comfort.
- The **HME Signature Series Dove Micro Low Air Loss Mattress System** may be used in a variety of settings including, but not limited to an individual in a home care setting or a long-term care setting who is suffering from skin breakdown, or pain management as prescribed by a physician.



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 **HME Signature Series Dove Micro Low Air Loss Mattress System** is an alternating pressure mattress replacement system used in the prevention and relief for patients with, or vulnerable to, pressure wounds. The mattress 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 **HME Signature Series Dove Micro Low Air Loss Mattress System** features a digital pressure adjustment function, mode selections, and audiovisual alarms. The 18 air cells in the mattress provide a unique design which keeps the lower sections of air cells constantly inflated while alternating and deflating the upper sections. The 3 head cells remain static and provide a "pillow" support for optimum comfort. The mattress has a heavy-duty polyester base sheet with a vapour-permeable two-way stretch with PU coated cover.

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



4. Product Installation Guide

Step 1 Step 2	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. Place the mattress on top of the bed frame. The feet symbol on both sides of the mattress indicates the location of the foot end.	HEAD END Air Mattress Control Unit CPR Valve Connector FOOT END
Step 3	Secure the mattress onto the bed frame by using the elastic straps or binding side straps	Securing Side Strap
Step 4	Ensure the CPR valve is at the CLOSE position before turning on the power.	
Step 5	Position the Control Unit by its elastic hanger brackets over the footboard of the bed. The elastic hanger brackets will self-adjust onto the footboard tightly	
Step 6	Remove the Transport Cap of the hose connector and connect the hose connector to the Control Unit. Firmly push the hose connector into position and a "click" sound will secure the connection.	

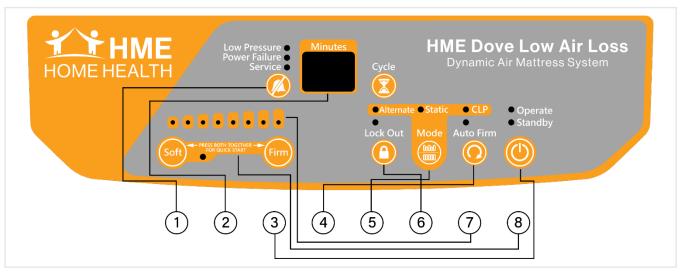


Direction for Connection Connect the power cord to the Pump. The power switch should remain off. Press the red power cord protector downward to secure the cord. Plug the power cord into the electrical outlet. NOTE: Check and ensure the Control Unit is suitable for the local power voltage. NOTE: When you first turn on the pump, allow Inflation Mode to complete before you select the Mode. Auto-Firm will flash while Inflation Mode completes. Once the pump is finished Auto-Inflation mode, you will see a dot in the bottom right corner of the display. You can then choose your desired Mode. CAUTION: The Control Unit can only be used with the mattress recommended by the manufacturer. Do not use the Control Unit for any other purpose. WARNING: Do not place the Control Unit in any area where the power cord can come off easily. For patient transportation, press the "Auto Firm" button and wait for 5 minutes for the mattress to be inflated. Step 7 Disconnect the hose from the Control Unit and put on the hose connector Transport Cap to keep the mattress inflated.



5. Panel Display & Operational Guide

5.1 Panel Display



	Alarm Mute & Alarm Indicator		
1	Low-Pressure Alarm Indicator		
*	Power Failure Alarm Indicator		
	Service (Malfunction) Alarm Indicator		
2	Alternating Cycle Time or Warning Code Display		
3	Operating or Standby		
4	Auto-Firm		
5	Functional Mode Selection (Alternate/Static/Constant Low Pressure)		
6	Panel Lock-out		
7	Comfort Control		
8	Auto Detection		

Alarm Mute Press the alarm mute button to temporarily suspend alarms. Should the situation not be resolved, and fault conditions continue, the alarm will resume notifying the patient and caregiver.
Alternate Time Display Alternate cycle time can be selected from 10-30 minutes by pressing the Cycle button.



	,			
	Operate or Standby Press this button to start operating or go into standby.			
	Auto-Firm The Control Unit will go into the inflation mode (LED lights flashing) every time the operate mode is triggered. This ensures the mattress can reach its maximum operating pressure. Once the max pressure level is reached, the Control Unit will automatically switch into the previously selected mode and comfort level. Users can also use this function as full mattress inflation during patient sit-ups or ingress/egress for better support.			
● Alternate ● Static ● CLP Mode	 Function Mode Switch Alternate – one of every three air cells slowly inflate and deflate to reduce surface pressure. Static-mattress cells maintain at the selected pressure. Automatically goes back to alternate mode after 20 minutes. Constant Low Pressure (CLP) – mattress cells maintain at the selected pressure and cycle time until another mode is selected. 			
	Panel Lock Should the panel remain untouched for 30 seconds, the Lock feature will automatically lock the panel to prevent anyone from accidentally changing settings without notice. To unlock, press the Lock button for 3 seconds.			
Soft Firm	Comfort Level Comfort level controls the air pressure output level. Press the Firm button and the output pressure will increase and higher-pressure output will support a higher weight; for decreasing air pressure, vice versa. Refer to Table 1 Weight and Comfort Level Reference for weight and comfort level suggestions.			
Soft PRESS BOTH TOGETHER - FIRM	Auto Detection Pressing both the Soft and Firm buttons together will activate Auto-Detection and Auto-Detection will automatically set the appropriate pressure output for the patient. When activated, the Auto-Detection light will flash to indicate the Control Unit is detecting the comfort level of the individual. Once the Control Unit has completed detecting, the Auto-Detection light will stop flashing and remain ON. The mattress pressure can be manually adjusted by pressing the Soft or Firm button if the individual wishes to change the comfort level.			



5.2 Operating Guide

5.2.1 General Operation



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

- Press "I" to turn on the unity, all LED indicators on the control panel will light up accompanied by a beep for 2 seconds (check for indicator failure if any), and the Standby indicator on the control panel will light up. If the Control Unit was previously shut off in operate mode then the Control Unit will enter operate mode directly.
- Press "O" to test the alarm battery this will turn off the power and the power failure should be triggered. (Refer to **Audio Visual Alarm** if the alarm is not triggered)
- Press the **Operate/Standby** 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 "IF" will be triggered.



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

- After initial inflation is completed, press the Auto-Firm button to transfer the person/user onto the
 mattress. The mattress will turn into a static condition in around 5 minutes. Move the patient onto the
 mattress and press the 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". Users can determine an appropriate pressure by adjusting the Comfort Level. Please
 consult with your physician for a proper setting.



Warning: The Control Unit should always be operating to prevent pressure wounds.

• In operating mode, press the **Operate/Standby** button for the system to enter standby mode. The system should be in standby mode before shutting down. Switch the **Power Switch** to off and the warning code "<u>Sd</u>" will appear on the display to shut off the system.



Note: A power failure alarm will be triggered if the power is switched off in operating mode (refer to Audiovisual Alarm). Press the power switch to restart the system or press Alarm Mute to turn off the system (refer to Alarm Mute).



Table 1: Weight & Comfort Level Reference

Comfort	Patient Weight								
Control	44 lbs.	88 lbs.	132 lbs.	176 lbs.	220 lbs.	264 lbs.	308 lbs.	352 lbs.	397 lbs.
(Indicators)	20 kg	40 kg	60 kg	80 kg	100 kg	120 kg	140 kg	160 kg	180 kg
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5.2.2 CPR

• When CPR needs to be performed, quickly rotate the CPR valve to the "OPEN" position, at the same time, disconnect the hose connector from the Control Unit to speed up the air release.

5.2.3 Audiovisual Alarm

 Power Failure – When electrical shortages occur or the power cord is unplugged without turning off the Control Unit or the Power Switch is pressed (intentionally or unintentionally), the "Power Failure" indicator will light up along with the buzzer and will last 20 minutes.



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

- Low Pressure When an abnormally low pressure occurs in the body section, the "Low Pressure" indicator will flash and beep. Should the situation not resolve and fault conditions continue, the alarm will resume.
- Service (Malfunction) When fault conditions occur, the "Service" indicator will light up along with a buzzer sound.

5.24 Alarm Mute

- When alarms are triggered, both the 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 be 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 the "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 the priority level. Refer to **Table 2 Warning Code Reference** for priority level.



Table 2: Warning Code Reference

Priority High to 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	S. d.	Power Failure	ONCE	POWER-OFF	System Shutdown	<u>S</u> hut <u>d</u> own
2	8 8	ALL LED	ONCE	OPERATE or STANDBY	Power-On	All Indicator Lights On
3	N/A	N/A	ONCE	OPERATE or STANDBY	State/Mode Switching	No Display
4	Ι. Ε.	Auto Firm	ONCE	OPERATE	Mattress Inflation Completion	<u>I</u> nflation <u>E</u> nded
5	R E.	Auto Firm	ONCE	OPERATE	Auto-Firm Completion	<u>A</u> uto Firm <u>E</u> nded
6	S. E.	Static	ONCE	OPERATE	Static Completion	Static Ended
7	N/A	Power Failure	REPEAT (Cycle 4 sec.)	POWER-OFF	Power Failure Alarm	No Display
8	I. F.	Low Pressure	REPEAT (Cycle 4 sec.)	OPERATE or STANDBY	Power-On Inflation Failure Alarm	<u>I</u> nflate <u>F</u> ailure
9	R F.	Low Pressure	REPEAT (Cycle 4 sec.)	OPERATE or STANDBY	Auto-Firm Failure Alarm	<u>A</u> uto-Firm <u>F</u> ailure
10	L, P,	Low Pressure	REPEAT (Cycle 4 sec.)	OPERATE or STANDBY	Low Pressure Overtime Alarm	<u>L</u> ow <u>P</u> ressure
11	H. P.	Service	REPEAT (Cycle 4.5 sec.)	OPERATE or STANDBY	High Pressure Overtime Alarm	<u>H</u> igh <u>P</u> ressure
12	H. F.	Service	REPEAT (Cycle 4.5 sec.)	OPERATE or STANDBY	High Ambient Temperature Alarm	<u>H</u> igh <u>T</u> emperature
13	U. I	Service	REPEAT (Cycle 4.5 sec.)	OPERATE or STANDBY	Air Valve 1 Positioning Failure Alarm	Air Valve 1 Failure
14	L. b.	Service	REPEAT (Cycle 15 sec.)	OPERATE or STANDBY	Battery Low Alarm	Battery may need recharging or may need to be replaced
15	C, U,	NONE	NONE	FACTORY CALIBRATION MODE	Calibration Not Completed	<u>C</u> alibration <u>U</u> nfinished
16		NONE	NONE	FACTORY CALIBRATION MODE	Calibration Completed	<u>C</u> alibration <u>C</u> ompleted



6. Cleaning

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



CAUTION: Do not immerse or soak the Control 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 placing it on 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, the mattress cover should be completely removed. Mattress covers can be laundered as follows: Prewash 60°C +15 minutes Main wash 60°C +15 minutes This should be followed by a cold rinse and extraction. 			
Drying	Mattress covers should be hung from a line or bar and drip dried in a clean indoor environment. The covers must be completely dried before returning to the mattress. Mattress covers can be tumbled dried on a low heat setting for 90 minutes. The drying temperature must not exceed 40°C. Exceeding the temperature can cause significant damage to the mattress cover.			



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



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



7. Storage

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

8. Maintenance

General	 Check the main power cord and plug for any abrasions or excessive wear. Check the mattress cover for signs of wear or damage. Ensure the mattress cover and tubes are stubbed together correctly. Check the air hoses for any kinks or breaks. For replacement, please contact your local dealer.
Fuse Replacement	 Disconnect the plug from the main 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 and replace the cover of the fuse holder. 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 mild detergent and water. Dry the filter before use. Check and replace the air filter regularly if the environment is dirty.

9. Disposal of Air Mattress

• When the air mattress is no longer useable, the mattress and the Control Unit may be discarded.



10. Troubleshooting

The mattress is not able to connect to the Control Unit:

- Check if the mattress model (model no. located inside the cover by the foot end) xxAAAxxx matches with the Control Unit model xxBBB-xxx. The AAA should be the same as BBB. If not, contact your dealer.
- Check if the Transport Cap is removed and make sure the connector is not broken.

The Control Unit is not working:

- Check if the plug is connected to the main supply.
- Check if the power switch is switched to the ON position.
- Check if there is a blown fuse.

Power failure alarm failed:

• If the Control Unit is in operation but failed to trigger the power failure alarm during power off. Charge the Control Unit for 6 hours or more of operating time and if the power failure still does not work, contact your dealer.

The low-pressure light is constantly flashing, and the alarm is sounding:

- Check if the CPR is in the CLOSE position
- Check if the connection between the air tubes to the Control Unit is tightly secured.
- Check if all coupling connections between the air cells and side rail are secured.
- If the main power supply is normal but there is no sound from the Control Unit, please remove the connector from the Control Unit to check if there is air coming out. If not, please turn off the Control Unit and contact your dealer.
- If all of the above steps have been checked. Press "Alarm Mute" for the system to be verified again.

The Control Unit is on, but the mattress is not alternating:

- Ensure the mattress has completely inflated.
- Check the control panel, the indicator light for "Alternate" should be on, if not, switch it to "Alternate."
- Check if the "Service" alarm indicator is on with the buzzer, if yes, contact the dealer.

Service (Malfunction) Alarm is on:

• Press "alarm mute" for the system to be verified again. If the alarm is still on, please contact the dealer.

The Control Unit is noisy:

- Make sure the Control Unit is resting against a solid surface.
- If the noise is getting louder, contact your dealer for further investigation.

The patient is bottoming out (without alarm triggered):

- Pressure settings 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 has sounded" for inspection.



11. Technical Data

11.1 Product Specification

Contr	ol Unit	Air Mattress		
MODEL	UD311-113 UD311-313 UD331-113 UD331-313	MODEL	81-311-002 81-331-004	
DIMENSION (cm)	33 (W) x 24 (D) x13 (H)	DIMENSION (cm)	200 (L) x 90 (W) x 20 (H)	
WEIGHT (kg)	3.5	WEIGHT (kg)	9	
CYCLE TIME	10/15/20/25/30 minutes	CELL MATERIAL	TPU	
STATIC TIME	20 minutes	NO. OF AIR CELL	18 Cells	
AUTO FIRM TIME	20 minutes	COVER MATERIAL	Two-way stretch polyester	
CONTROL UNIT OUTPUT FLOW RANGE (Litre)	> 8L (120V) Note: The flow rate may be varied because of the fluctuation of input voltage	BOTTOM MATERIAL	Polyester-PU	
CONTROL UNIT OUTPUT PRESSURE RANGE (mmHg)	15 to 50 (±5)	MAX WEIGHT	350 lbs	
POWER	AC120 V 60Hz			
CURRENT	0.25AMAX (@132V~)			
FUSE RATING	T1A 250VAC			
FREQUENCY	60 Hz (120V)			
CLASSIFICATION	Class II Type BF			
WARRANTY	2 years	WARRANTY	2 years	
OPERATION ENVIRONMEN	ІТ	5°C ~40°C 15%RH ~ 93%RH (no condensation)		
STORAGE ENVIRONMENT		-25°C ~70°C ≦ 93%RH (no condensatio	n)	
ENVIRONMENT PRESSURE		70 kPa-101.3kPa		
ENVIRONMENT HORIZON		≦3000m		
WATER AND DUST PROTEC	CTION CLASSIFICATION	IP21		



11.2 EMC Information (120V)

Guidance & Manufacturer's Declaration-Electromagnetic Emissions					
The <u>device(s)</u> is intended for use in the electromagnetic environment specified below. The customer or the user of the <u>device(s)</u> should assure that it is used in such an environment.					
Emission test	Compliance	Electromagnetic Environment-Guidance			
RF emissions CISPR 11	Group 1	The <u>device(s)</u> 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 <u>device(s)</u> is suitable for use in all establishments, including domestic			
Harmonic emissions IEC 61000-3-2	Class A	establishments and those directly connected to the public low-voltage power supply network			
Voltage fluctuations/flicker emissions IEC 61000-3-3	Compliance	that supplies buildings used for domestic purposes.			



Guidance & Manufacturer's Declaration-Electromagnetic Immunity

The <u>device(s)</u> is intended for use in the electromagnetic environment specified below. The customer or the user of the <u>device(s)</u> should assure that it is used in such an environment.

Immunity Test	IEC 60601 Test Level	Compliance Level	Electromagnetic Environment-	
minumey rest		Compliance Level	Guidance	
Electrostatic	+ 6 kV contact	+ 6 kV contact	Floors should be wood, concrete, or	
discharge (ESD) IEC	+ 8 kV air	+ 8 kV air	ceramic tile. If floors are covered with	
61000-4-2			synthetic material, the relative humidity	
01000-4-2			should be at least 30%	
Electrical fast	+ 2kV for power	+ 2kV for power	Mains power quality should be that	
	supply lines	supply lines	of a typical commercial or hospital	
transient/burst IEC 61000-4-4	+ 1kV for	Not applicable	environment.	
01000-4-4	input/output lines			
Surge IEC 61000-4-5	+ 1kV line(s) to	+ 1kV differential	Mains power quality should be that	
	line(s)	mode Not applicable	of a typical commercial or hospital	
	+ 2kV line(s) to earth		environment.	
	<5% UT (>95% dip in	<5% UT (>95% dip in	Mains power quality should be that	
Voltage Dips, short	UT) for 0,5 cycle 40%	UT) for 0,5 cycle 40%	of a typical commercial or hospital	
interruptions, and	UT (60% dip in UT)	UT (60% dip in UT)	environment. If the user of the	
voltage variations on	for 5 cycles	for 5 cycles	device(s) requires continued operation	
power supply input	70% UT (30% dip in	70% UT (30% dip in	during power mains interruptions, it is	
lines IEC 61000-4-11	UT) for 25 cycles	UT) for 25 cycles	recommended that the device(s)	
illes IEC 61000-4-11	<5% UT (>95% dip in	<5% UT (>95% dip in	be powered by an uninterruptible	
	UT) for 5 s	UT) for 5 s	power supply or a battery.	
	3 A/m	3 A/m	The <u>device(s)</u> power frequency	
Power frequency			magnetic fields should be at levels	
(50/60 Hz) magnetic			characteristic of a typical location.	
field IEC 61000-4-8			in a typical commercial or hospital	
			environment.	
NOTE: UT is the a.c. m	ains voltage prior to app	olication of the test leve	l.	



Guidance & Manufacturer's Declaration-Electromagnetic Immunity

The <u>device(s)</u> is intended for use in the electromagnetic environment specified below. The customer or the user of the <u>device(s)</u> should assure that is used in such and environment.

Immunity Test	IEC 60601 Test	Compliance Level	Electromagnetic Environment-
Conducted RF IEC 61000-4-6 Radiated RF IEC 61000-4-3	3 Vrms 150 KHz to 80 MHz 3 V/m 80MHz to 2,5 GHz	3 Vrms 3 V/m	Portable and mobile RF communications equipment should be used no closer to any part of the device(s) including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter. Recommended separation distance: $d = 1,2 \sqrt{P}$ $d = 1,2 \sqrt{P}$ 80 MHz to 800 MHz $d = 1,2 \sqrt{P}$ 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 distance in metres (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: ((a))

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.

- 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 device(s) is used exceeds the applicable RF compliance level above, the device(s) should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the device(s).
- b. Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3 V/m.



Recommended Separation Distance Between Portable & Mobile RF Communications Equipment and The <u>Device(s)</u>

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

Rated maximum output power of transmitter	Separation Distance According to Frequency of Transmitter m			
	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2,5 GHz	
W	$d = 1,2 \sqrt{P}$	$d = 1,2 \sqrt{P}$	$d = 2,3 \sqrt{P}$	
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 metres (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.	



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