

# APH052 Apollo 5 Plus<sup>™</sup> APH053 Apollo 8<sup>™</sup>



# **USER MANUAL**

AHSM-001 Version 5 08.4.2021

www.apollo-ht.co.uk

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## INSTRUCTIONS

#### **DANGER** - To reduce the risk of electrocution:

- 1. Do not place or drop into water or liquid.
- 2. Do not reach for a product that has fallen into water.
- 3. Do not place or store the product where it may fall into water.
- 4. Always unplug this product when not in use.

#### WARNING - To reduce the risk of burns, electrocution, fire or injury to Persons:

- 1. Do not block air openings of this unit or place it on a soft surface.
- Do not operate the unit if it has a damaged cord or plug, return the unit or contact a service centre for examination or repair.
- 3. Do not drop or insert any object into any opening or hose.
- 4. Connect the unit to a properly grounded outlet only. See grounding instruction.
- 5. When plugged in, the unit should not be left unattended.
- 6. The unit is only for it's intended use, do not use attachments not recommended by the Manufacturer.
- 7. Close supervision is necessary when this product is used near Children.
- 8. Keep the cord away from heated surfaces.

#### **NOTE, CAUTION AND WARNING STATEMENTS:**

**CAUTION** – Indicate correct operating or maintenance procedures in order to prevent damage to or destruction of the equipment or other property. **WARNING** – Calls attention to a potential danger that requires correct procedures or practices in order to prevent personal injury.

# **SYMBOLS**

<u> </u>	Caution
	Manufacturer
★	Complies with standards protecting against electric shock for type BF equipment.
Ţ <u>i</u>	Consult operating instructions for use
IP21	Protected against solid foreign objects of 12.5 mm and greater; Protection against vertically falling water drops
	Class II
	Temperature limitation/temperature range
P	Dry clean, Any Solvent Except Trichloroethylene
A Company	Do Not Iron
$\odot$	Tumble Dry, Normal, Low Heat
X	Do Not Tumble Dry
×	Do Not Bleach
$\boxtimes$	Do Not Dry Clean
95	Machine wash, regular / normal, max 95 degrees C (203 degrees F)
X	Attention – Observe proper Disposal of Electrical & Electronic Equipment (WEEE): This product should be handed over to an appropriate collection point for the recycling of electrical and electronic equipment. For more detailed information about the recycling of this product, please contact your local city office, household waste disposal service or the retail store where you purchased this product.

# A. INTRODUCTION

For initial set up and for reference purpose, this manual should be read carefully.

#### **A.1 GENERAL INFORMATION**

The unit is for prevention or treatment of pressure ulcers.

The unit has been tested and successfully approved by the following regulation:

IEC/EN 60601-1

IEC/EN 60601-1-2

IEC/EN 60601-1-11

IEC/EN 61000-3-2 Class A

IEC/EN 61000-3-3

CISPR 11 Group 1, Class B



#### **A.2 INTENDED USE**

The unit should help prevent and reduce the incidence of pressure ulcers while optimising Patient comfort. It is used for the following purposes:

- 1. For use in Hospitals, rehabilitation facilities, long term home care
- 2. Patients up to very high risk of developing pressure Ulcers
- 3. Patients who present with pressure ulcers which need treatment management
- 4. Patient who may suffer Pressure Area Care Damage.
- 5. To assist in the prevention of pressure area damage.

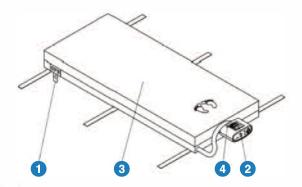
#### **A.3 CONTRAINIDITIONS**

The mattress is not suitable for use on patients with cervical or skeletal traction and unstable spinal cord injuries.

NOTE: The unit is not suitable for use around flammable anaesthetic mixture, with air or with nitrous oxide.

# **B. PRODUCT DESCRIPTION**

#### **B.1 AIR PUMP AND MATERIALS**



- 1 CPR
- 2 Pump unit
- 3 Mattress System
- 4 Quick Connectors

#### **B.2 PUMP AND MATERIALS**



- 1 Comfort Adjust button (30 200kg)
- 2 Power On/Off button
- 3 Mode button (Alternating/Static)
- 4 Alarm Mute button
- 5 Panel Unlock button
- 6 Fuse
- 7 Air Filter
- 8 Mounting Brackets
- 9 Power Cable Plug



#### **B. 3 FRONT PANEL**

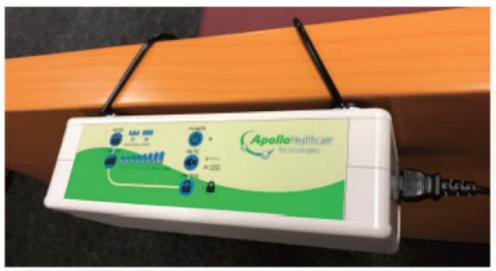


- Comfort (weight) setting indication lights. Adjust the comfort button through the weight settings from 30KG to 200KG, according to the patient's weight.
   The weight settings should only be used as a comfort setting and as a guide in conjunction with a qualified Clinicians advice for each individual user.
- 2. Fault indication light. If the pump detects low pressure in the mattress or the pump requires a service, this light will illuminate Orange, an audible constant beep alarm will also sound. Once the issue has been rectified press the Alarm Mute Button in order to return the pump to normal operation.
- 3. Panel lock out light. After 2 minutes if no buttons have been pressed on the pump, the panel automatically locks-out. This light will illuminate to show that the panel has locked-out. To unlock the panel, hold down the panel unlock button for 2 seconds. The light goes out and an audible beep will sound.
- 4. Alternating/Static indicator lights. Depending on whether the alternating or static function has been selected, the corresponding light will illuminate. Pressing the mode button will switch between alternating and static. As a safety feature, the pump automatically switches to alternating from static after a period of 30 minutes.
- 5. Alarm mute button. When either the fault or power failure alarms are triggered, pressing this button will temporarily silence the alarm whilst the issue is fixed.
- 6. Power failure alarm indicator light. If the power is lost to the pump this light will illuminate Red and a constant beep alarm will sound. Once the power is restored to the pump the light and alarm will stop.
- 7. Power on/off indicator light. When the pump is turned on by pressing the power button, this light will illuminate green. When the button is pressed again to turn off the pump or if there is a power failure, the light will go out.

## C. INSTALLATION

After delivery check the system for any damage which may have occurred during transit. If damaged in any way contact your supplier immediately.

#### C1 PUMP AND MATTRESS INSTALLATION



- 1. Place air mattress on top of the bed frame/platform. Please ensure foot sign is at the bottom of the bed.
- 2. Hang the pump onto the foot end bed rail and adjust the hangers to a suitable position for the pump. Alternatively, the pump may be placed on a flat surface.
- 3. If the pump wire mains wire could be a risk on the floor under the bed with the use of other equipment, clip the mains cable into the cable management system on the side of the mattress
- 4. Connect the air hose connectors from the mattress to the pump unit. Make sure the connection is completely secured.



NOTE: Ensure the air hoses are not kinked or tucked under the mattress.

5. Plug power cord into electrical socket.



NOTE: 1. Make sure the pump is suitable for the power voltage

2. the plug will disconnect the system



**CAUTION:** The pump can only be used for the mattress recommended by the Manufacturer. Do not use for any other purpose.

6. Press the power button to turn the pump on.

# D. OPERATION



NOTE: Always read the manual before use.

#### **D.1 GENERAL OPERATION**

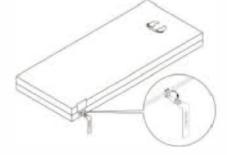


- 1. Insert the supplied power cable into the side of the pump and the plug into a suitable power supply and press the power on button.
- 2. Wait for 20-30 minutes (depending on the mattress) for the mattress to be fully inflated by the pump.
- 3. The pump will automatically be set to alternating mode and on the lowest comfort setting. Static mode can be selected if desired and the comfort setting can be increased. The alternating or static light will continue to flash until the mattress reaches the pressure chosen.
- 4. Adjust the comfort setting to the most comfortable level for the Patient, without the mattress bottoming out. The pressure in the mattress will slowly increase or decrease to the required level after the alternating or static light has stopped flashing.

NOTE: It is recommended that the comfort setting be set to maximum(200KG) for the guickest inflation of the mattress for every first inflation or new use, the user can then adjust the comfort setting to the desired level after the mattress has fully inflated.

#### D.2 CPR

This is for emergency use to deflate air from the mattress. Disconnect the air tubes from the mattress for even faster deflation.



#### **D.3 PRESSURE SET UP**

Press the comfort button to cycle through the different weight settings until the green LED on the desired weight is illuminated.

The alternating or static LED will flash until the pressure in the mattress reaches the desired setting.

#### **D.4 LOW PRESSURE INDICATION FUNCTION**

Once the mattress has reached the desired comfort setting, should the pump detect low pressure in the mattress the fault LED will light up Orange and an audible constant beep will sound. Check that the connector is fully inserted and that the CPR valve on the mattress is fully closed.



NOTE: If the pressure remains at LOW PRESSURE, check for

any air leakage (cells, air tubes and connectors) If necessary, replace any damaged parts. Or contact the supplier for further assistance.

#### **D.5 STATIC MODE**

If required, the mode button can be pressed to place the mattress into static. The pressure inside the cells will be inflated to the same comfort setting and the alternating function will be suspended. Pressing the mode button again will return the mattress back to alternating mode. As a safety feature the pump will AUTOMATICALLY switch back to alternating after a period of 30 minutes in static mode.

#### **D.6 ALARM MUTE**

When low pressure is detected due to a fault both alarm LED will light and audible alarm will sound. By pressing the alarm mute button, this will temporarily mute the audible alarm until the fault is rectified. Should the fault remain unresolved after 5 minutes, the alarm will sound again, the alarm LED light will remain on until the problem is solved. Contact your equipment provider or technician should the problem persist.

## E. CLEANING

Ensure the system is thoroughly cleaned between Patient use. Wipe the air pump with a damp cloth pre-soaked with a mild detergent and keep it dust free.



CAUTION, do NOT soak the pump.

Wipe the mattress with a damp cloth pre-soaked with warm water (not to exceed 60° C) containing a mild detergent. Keep the Mattress dust free. The cover may also be cleaned using 1000ppm Sodium Hypochlorite diluted in water. All parts should be air dried thoroughly before use. The mattress cover may also be washed in the washing machine at a temperature not exceeding 95° C. See printed washing instructions on the cover.



## **DO NOT USE PHENOLS**



Always dry the Mattress without direct exposure to sunlight.

## F. STORAGE

- 1. For storage fully deflate the mattress.
- 2. Roll the Mattress from head end downwards.
- 3. Store in a cool dry place.



NOTE: DO NOT FOLD OR STACK THE MATTRESS.

# G. MAINTENANCE

#### **G.1 GENERAL**

- 1. Check the main power cord and plug are not excessively worn or cut.
- 2. Make sure mattress cover, inner cells and air tubes are assembled correctly.
- 3. Make sure all the cells are alternating correctly.
- 4. Check all tubing is not kinked or broken. If replacements are required, contact your supplier.

#### **G.2 FUSE REPLACEMENT**

- 1. Firstly unplug the pump from the mains.
- 2. Remove the cover of the fuse with a cross head screwdriver.
- 3. Insert a new fuse of the correct rating and replace the cover. The fuse should be rated as T1A/250V and CE approved.

#### **G.3 AIR FILTER REPLACEMENT**



- 1. Remove the air filter cover, located at the back of the pump.
- 2. Replace the filter and replace the cover.
- 3. Replacing the air filter regularly is recommended.

# H. SPECIFICATIONS

Item		Specification	
Power Supply (Note: Seerating label on the product)		AC 220-240V 50 Hz, 0.09A (for 230V system)	
Fuse Rating		500mA, 250V	
Cycle time		12mins	
Dimension (L x W x H	)	27.5x13.5x9cm	
Weight		1.3Kg	
	AtmosphericPressure	80KPa to 106KPa	
		Operation: 5°C to 35°C (41°F to 95°F)	
	Temperature	Storage: -15°C to 50°C (5°F to 122°F)	
Environment		Shipping: -15°C to 70°C (5°F to 158°F)	
		Operation: 10% to 90% non-condensing	
	Humidity	Storage: 10% to 90% non-condensing	
		Shipping: 10 % to 90% non-condensing	
Classification		Class II, Type BF, IP21	
		Applied Part: Air Mattress	
		Not suitable for use in the presence of a flammable	
		anesthetic mixture (No AP or APG protection)	

Air mattress	Specification		
Mattress Height	APH052 - 7" or 17.5cm	APH053 - 8" or 20cm 200 x 90 x 20cm	
Dimension (L x W x H)	200 x 90 x 17.5cm		
Mattress Weight	9 kg (u with fo am base)	9 kg	
System Pressure Range	30 - 80 mmHg	30 – 80 mmHg	
Maximum load	200 kg	200 kg	
Suggested load	1st LED: 30~40 kg 2nd LED: 50~60 kg 3nd LED: 70~80 kg 4th LED: 90~110 kg 5th LED: 120~140 kg 6th LED: 150~170 kg 7th LED: 180~190 kg 8th LED: 200 kg	1 <sup>st</sup> LED: 30~40 kg 2 <sup>nd</sup> LED: 50~60 kg 3 <sup>rd</sup> LED: 70~80 kg 4 <sup>th</sup> LED: 90~110 kg 5 <sup>th</sup> LED: 120~140 kg 6 <sup>th</sup> LED: 150~170 kg 7 <sup>th</sup> LED: 180~190 kg 8 <sup>th</sup> LED: 200 kg	



NOTE: If necessary, please ask your supplier for further technical documents

# 10. Technical Specification:

Item			Specification		
Power Supply (Note: Seerating label on the product)		uct)	AC 220-240V 50 Hz, 0.09A (for 230V system)		
Fuse Rating			500mA, 250V		
Cycle time			12mins		
Dimension (L x W x H)			27.5x13.5x9cm		
Weight			1.3Kg		
	AtmosphericPressur	re	80KPa to 106KPa		
				Operation: 5°C to 35°C (41°F to 95°F)	
	Temperature		Storage: -15°C to 50°C (5°F to 122°F)		
Environment			Shipping: -15°C to 70°C (5°F to 158°F)		
	Humidity		Operation: 10% to 90% non-condensing		
			Storage: 10% to 90% non-condensing		
			Shipping: 10 % to 90% non-condensing		
Classification			Class II, Type BF, IP21		
			Applied Part: Air Mattress		
			Not suitable for use in the presence of a flammable		
			anesthetic mixture (No AP or APG protection)		
Mattress			Specification		
Model	T01	T05		T04(Cell-on-cell mattress)	
Dimension (L x W x H)	200x90x10cm	200x9	90x10cm	200x90x20cm	
Weight	5.31Kg	4.58Kg		9Kg	

180Kg

200Kg



# NOTE-

Max. Support Weight

- 1. Consult the distributor or EU representative for further technical documents.
- 2. The specification is also suitable for other areas operating with same power supply.
- 3. Mattress dimension and weight is measured without foam cushion.

180Kg

4. The manufacturer reserves the right to modify the specification without notice.

#### Appendix A: EMC Information

Guidance and manufacturer's declaration-electromagnetic emissions		
The device is intended for use in the electromagnetic environment specified below. The customer or the user of the device		
should assure that they are used in such an environment.		
<b>Emissions test</b>	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11	Group 1	The models device use RF energy only for their internal function. Therefore, their RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR11	Class B	The model P08E are suitable for used in domestic stablishment and in establishment directly connected to a low voltage power supply network which supplies buildings used for
Harmonic emissions IEC 61000-3-2	Class A	domestic purposes.
Voltage fluctuations / flicker emissions IEC 61000-3-3	Complies	

#### Warning:

- 1. The device should not be used adjacent to or stacked with other equipment. If adjacent or stacked use is necessary, the device should be observed to verify normal operation in the configuration in which it will be used.
- 2. Use of accessories, transducers and cables other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.
- 3. Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the Pump, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

#### **Guidance and Declaration-electromagnetic immunity**

The models device intended for use in the electromagnetic environment specified below. The customer or the user of the device should assure that they are used in such an environment.

Immunity test	IEC 60601	Compliance	Electromagnetic environment -guidance
,	test level	level	
Electrostatic	±8 kV contact	±8 kV contact	Floors should be wood, concrete orceramic tile. If
discharge	±2 kV, ±4 kV, ±8 kV,	±2 kV, ±4 kV, ±8kV, ±15 kV	floors are coveredwith synthetic material, the
(ESD) IEC	±15 kV air	air	relativehumidity should be at least 30 %.
61000-4-2			
Electrical fast	±2kV for power supplylines	±2kV for powersupply lines	Mains power quality should be thatof a typical
transient/burs	±1 kV for Input/outputlines		commercial or hospitalenvironment.
t			
IEC 61000-4-4			
Voltage dips,	<5 % UT	<5 % UT	Mains power quality should be thatof a typical
short	(>95% dip in UT.)	(>95% dip in UT.)	commercial or hospitalenvironment. If the user of
interruptions	for 0.5 cycle	for 0.5 cycle	themodel P08 require continuedoperation during
and	<5 % UT	<5 % UT	power mainsinterruptions, it is recommended that the
voltage	(>95% dip in UT)	(>95% dip in UT)	model P08 poweredfrom an uninterruptible
variations on	for 1 cycle	for 1 cycle	powersupply or a battery.
power supply	70% UT	70% UT	
input	(30% dip in UT)	(30% dip in UT)	
lines	for 25/30 cycles	for 25/30 cycles	
IEC	<5% UT	<5% UT	
61000-4-11.	(>95 % dip in UT)	(>95 % dip in UT)	
D	for 5/6 sec	for 5/6 sec	David francisco de la contra dela contra de la contra dela contra de la contra del la contra de la contra de la contra del la
Power	30 A/m	30 A/m	Power frequency magnetic fields should be at levels
frequency (50/60 Hz)			characteristic ofa typical location in a typicalcommercial or hospitalenvironment.
magnetic field			typicalcommercial of hospitalentificinitent.
IEC 61000-4-8			
	I. a.c. mains voltage prior to application (	of the test level	
Conducted	3 Vrms	3 Vrms	Portable and mobile RF communicationsequipment
RFIEC	150 kHz to 80 MHz	150 kHz to 80 MHz	
61000-4-6	6 Vrms in ISM andamateur	6 Vrms in ISM andamateur	should be used no closer to any part ofthe models
	radiobands	radiobands	P08 including cables, than therecommended
			separation distance calculated fromthe equation
			applicable to the frequency of thetransmitter.
			Recommended separation distance
			$d=[3,5/V1]x P^{1/2}$
Radiated	10 V/m	10 V/m	d=1,2xP <sup>1/2</sup> 80 MHz to 800 MHz
RFIEC	80 MHz to 2.7 GHz.	80 MHz to 2.7 GHz	d=2,3xP <sup>1/2</sup> 800 MHz to 2.7 GHz
61000-4-3			u-2,5XP 800 WITZ to 2.7 GHZ
	385MHz-5785MHz	385MHz-5785MHz	where P is the maximum output power rating of
	TestspecificationsforENCLOSUREP	Testspecificationsfor	thetransmitter In watts (W) according to the
	ORTIMMUNITY toRF	ENCLOSUREPORTIMMUNIT	· · ·
	wirelesscommunicationequipmen	Y toRF	transmittermanufacturer and d Is the
	t(Refer to table9 of IEC	wirelesscommunicationeq	recommendedseparation distance in meters (m).
	60601-1-2:2014)	uipment(Refer to table9 of	Field strengths from fixed RF transmitters,
		IEC 60601-1-2:2014)	asdetermined by an electromagnetic site
			survey, ashould be less than the compliance level in
			eachfrequency range. Interference may occur in the
			vicinity of equipmentmarked with the following
			· · · · ·
	İ	Í	symbol:
			l I
			// \
			(((•)))
			$((\bullet))$
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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 and 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 models P06A is used exceeds the applicable RF compliance level above, the model P06A should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the model P06A.
b) Over the frequency range 150 kHz to 80 MHz, field strengths should be less than 3V/m.

#### Recommended separation distances between portable and mobile RF communications equipment and the model P08E

The model P08E is intended for use in electromagnetic environment in which radiated RF disturbances is controlled. The customer or the user of the model P08E can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the model P08E is recommended below, according to the maximum output power of the communications equipment.

Rated maximum outputPowerof transmitter		Separation distance according to frequency of transmitter m	
w	<b>150kHz to</b> <b>80MHz</b> d=1.2xP <sup>1/2</sup>	80MHz to 800MHz d=1.2xP <sup>1/2</sup>	<b>800MHz to 2,5GHz</b> d=2.3xP <sup>1/2</sup>
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) accord able 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.

# SAVE THESE INSTRUCTIONS

It is **NOT** recommended to repair or replace the unit, unless this is by qualified personal. To reduce the risk of shock or electrocution do not change or modify the cord or plug in any way.



For further technical support please visit www.apollo-ht.co.uk





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Due to ongoing research and development, Apollo Healthcare Technologies Ltd, reserve the right to change specifications without prior notice. This will not affect the efficacy of the system. Always consult the user manual for instructions for use.