

APH055 Apollo 4 Plus[™]



USER MANUAL

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www.apollo-ht.co.uk

INTRODUCTION

Congratulations on the purchase of your Apollo 4 Plus alternating pressure cushion. The durable, high quality material used in the manufacturing of these products will ensure that you experience long-lasting and uninterrupted performance.

Intended Use

This product is intended to help and reduce the incidence of pressure ulcers while optimising patient comfort. It also provides the following benefits:

- To help and reduce the incidence of pressure ulcers while optimising patient comfort.
- For long term home care of patients suffering from pressure ulcers.
- For pain management as prescribed by a physician.

The product can only be operated by personnel who are qualified to perform general nursing procedures and have received adequate training in knowledge of prevention and treatment of pressure ulcers.

Contraindication

Patient conditions for which the application of pressure relieving therapy on an alternating system are contraindicated as follows:

- Cervical or skeletal traction
- Unstable spinal cord injuries



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Table of Contents

Symbol definition
Products Feature
Safety Precaution
Confirm before fixing
Parts name
User instructions
Maintenance
Storage and disposal methods
Troubleshooting
Warranty and after-sales service
Specification list
EMC Declaration

Symbol definition



CE Mark

EC REP

European representative information

LOT

Product production batch

SN

Product serial number



Manufacturer information



II type equipment symbol



Mandatory reading of instructions



BF device symbol

IP21

Protected against solid foreign objects of 12.5mm and greater; protection against vertically falling water drops.



When the end user intends to discard this product,

the product must be sent to the appropriate facility for recycling and recycling.



Environment temperature for operation or storage



Environment humidity for operation or storage



Atmospheric pressure for operation or storage



Maximum weight of the cushion load.



Do not wash the cushion.



Do not bleach the cushion.



Do not tumble dry.



Do not iron.



Do not dry clean.

Products Features

1. The product consists of an alternating pressure air pump powered by an internal lithium battery and a moulded visco foam cushion base with 6 alternating cells. It is generally installed on a wheelchair and used as a cushion.





Safety Precautions

In order to use this product safely, please read this IFU carefully and understand it. Please be sure to follow the instructions in this manual. In order to avoid a disaster when using this product, you must know in advance the factors that may be dangerous. However, it is difficult to predict all potential dangers. Therefore, warnings about security are defined and described in this manual.

Warning If you do not follow the instructions in this symbol, you are warned that there is a risk of property damage, injury or death. Because it is especially important, it is described below as "safety precautions and warnings"

Safety warnings

Warning 1 This product features a layer of soft air floating beneath the patient's body. Therefore, do not use when patients need to undergo cardiac resuscitation.

Warning 2 Use this product with your doctor or specialist when using this product. In addition, if the body feels abnormal during use, or the symptoms are worse, or there is a risk of accident, stop using it immediately and consult an expert.

Warning 3 When you move to a stand position or get out of cushion, be sure to ask someone else for assistance, otherwise there is a danger of falling.

Warning 4 Smoking is not allowed on the cushion and may cause a fire.

Warning 5 Please do not bundle the air outlet or press it under the cushion. When the blower tube is bent or compressed, air may not enter the air mattress and may not achieve the desired results.

NO SMOKING

Warning 6 Do not use the air pump in a damp place. Do not use in or around water or urine, which may result in electric shock or malfunction. Avoid direct sunlight. In hot and humid places, keep it away from the wall or above 55cm to avoid moisture retention.

Warning 7 Please do not remove the screw of the air pump to open the casing and repair it yourself; it may cause electric shock or malfunction. In addition, the modification of the air pump without our consent may pose a serious safety hazard. Never let customers modify the air pump.

Warning 8 If you do not use the air pump for a long time or need to clean the air pump, be sure to unplug the power cord from the outlet. If not it may cause accidents, electric shocks, and malfunctions.

Warning 9 Please do not pull on the power cord of the air pump, do not damage the power cord, and do not clip the power cord to the door. When removing the plug from the socket, be sure to hold the power plug and pull it out to avoid electrical accidents and fire.

Confirm before using

Before using, please confirm whether the accessories are complete and whether you understand the contents and precautions of the manual.

Packing list confirmation

1 Air pump for cushion	x1
2 Power cord	x1
3 Power Adapt	x1
4 User manual	x1
5 Cushion	v1







User instructions

Step1:

Place the cushion on the chair

N.B. Actual cushion may look different to that shown in the image



Step2:

- Hang the hooks of the air pump on the chair frame or put on a nearby platform if the chair is not moving
- Connect the air pipes using the quick connector to the side of the pump



N.B. Actual pump may look different to above image

Step3:

- Turn On the power 🕦
- Adjust the Pressure 2
- Adjust the Function (3)



- 1 Power: Press this button to turn the pump on and off.
- Pressure: Press this button to cycle through the 3 pressure settings: LOW = 20mmHg MEDIUM = 40mmHg HIGH = 60mmHg
- 3 Static: Press this button to cycle through the Static & Alternate settings: Default setting is 6 min alternating cycle. You can select to put the pump into static function for 10 min or 20 min. After the selected time has elapsed, the pump automatically defaults to 6 min alternating cycle
- 4 Battery Indicator: Shows the level of charge currently in the battery. When all 4 lights are illuminated green, the battery is fully charged. As power is used, the number of lights illuminated reduces. When there is only 1 light left, it is time to charge the battery.
- Charging Indicator: When the power adapter provided with the pump is connected to the pump and to a suitable power supply, the charging indicator symbol is illuminated green.

As the battery charges the lights on the battery symbol illuminate gradually from no lights to 4 lights and then all lights extinguish and the sequences starts again until the battery is fully charged.

When the battery is fully charged, all 4 lights on the battery symbol remain illuminated constantly.

Once the battery is fully charged remove the power adapter from the pump, although there is a safety feature to prevent overcharging.

Step4: Terminate operation of device

 When you do not need to use the device, turn off the power to save the charge in the battery.



Please use the adapter provided by the manufacturer to charge the battery, otherwise it may cause the battery affect the life of the battery, or cause the battery to overcharge, or cause fire and other hazards.

Maintenance

Warning: Do not carry out maintenance when the product is in use.

Air pump output pressure check

In the case of air pumps and air cushions, please check the pressure once a
week. In addition, if you change the location, or if you suspend operation
due to power outages, etc., please do it every time, even within a week.

Cushion cleaning

- Usually use a dry cloth to clean the cushion
- If necessary, wet the cloth with a neutral detergent, wring the cloth and clean the cushion cover.
- Let it dry naturally
- Maintenance is required when the product is no longer in use.

Maintenance is required when the cushion is contaminated by the patient's bodily fluids. If the inner foam becomes wet, it must be replaced.

Maintenance is required every other week when in use.

Warning:

- Dry cleaning is forbidden, otherwise it will hurt the cushion
- Please do not use a dryer to avoid deformation of the cushion.
- Do not use an iron to avoid damage to the cushion cover

Air pump cleaning

- Turn off the power switch and unplug the power cord.
- Usually use a dry cloth to clean the air pump.

Storage and disposal methods

If you are no longer using this product, please follow the steps below.

- 1. Turn off the power switch and unplug the power cord.
- 2. Unplug the air tube
- 3. Cleaning dirty air pumps and cushions
- 4. Do not fold the cushion and put it in the plastic bag
- 5. Tighten the power cord and put the air pump in a plastic bag.
- 6. Put the air pump and cushion in the box, together with this manual

Note:

- Please put it in a stable place, do not drop from a height.
- Please do not put heavy objects on the box.
- Please keep it in a place with low humidity.

Troubleshooting

Trouble	Possible reason
Pump does not start, no light on	The battery is flat or the power plug is
power button	not plugged into the power outlet,
	the power supply does not work.
Air cushion is too soft	Valve does not work
	Connector is off
	Air pipes kinked
	Damaged cell inside
	cushion
Air cushion is too Firm	Pressure Selection error
Air pump noise is abnormally	Put something on the air pump
high	The air pump is in contact with other
	things
	Put the air pump on something that is
	easy to vibrate
Power outage	The power plug
	Internal fault

Warranty and after-sales service(Please read carefully)

Service life and Shelf life

- EXPECTED SERVICE LIFE OF the pump is 5 years
- EXPECTED SERVICE LIFE OF the cushion is 2 years

Warranty

Item	Warranty
Pump	24 months
Cushion	12 months

About the repair request

First, look at the possible causes (see page 14). If you are still having trouble, turn off the product and contact your dealer or Apollo Healthcare Technologies.

During the warranty period:

- In the event of a malfunction in a normal use condition, your dealer should repair according to the provisions of the warranty.
- Please show your warranty when you are under warranty.
- In addition, if the second article of the guarantee is met, the repair is charged. Please confirm the guarantee for details.

After the warranty expires:

 Please consult with the shop or company you purchased, and if you can maintain the function through maintenance, you can repair it at a charge according to the customer's request.

Service personnel requirements

- Should have a certain understanding of the product.
- Should be able to understand the circuit schematic.
- Should be able to use screwdriver, soldering iron, and multimeter, etc.
- We (manufacturer) will provide the technical instructions, the circuit diagram, parts list to the service personnel.

Warning:

- Please don't replace the power cord by yourself and it must be replaced by professional service personnel.
- The parts should be replaced by professional service personnel when it is necessary to be replaced.
- The only safe way to turn off the power completely is to unplug the power cord

Specification list

Specifications

Adapt Power Supply		Input: AC 100-240Vac 50/60Hz 1.6A	
Adapti Gire. Supp.,			
		Output: 28Vdc 2.0A	
Adapt dimension	ı(LxWxH)	12x13.5x5.5 cm	
Internal Li-Batter	Ty .	22.2Vdc 2000mAH	
Work time whe	en the battery is fully	8 hours	
charged			
Cycle time		6 mins, Fixed, non-adjustable (*Customization)	
Dimension(LxWx	:H)	27.5x14x9.5 cm	
Weight		1.8Kg	
Environment	Atmospheric pressure	80KPa to 106KPa	
	Temperature	Operation: 5° C to 40° C (50° F to 104° F)	
		Storage: -15° C to 70° C (5° F to 158° F)	
		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/Air pad	
		Not suitable for use in the presence of a	
		flammable	
		Anesthetic mixture (No AP or APG protection)	
Cushion		Size 45x45x10cm	
		Weight:1.0Kg	

EMC Declaration

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.

Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions	Group 1	The models device use RF energy only for their internal
CISPR 11		function. Therefore, their RF emissions are very low and
		are not likely to cause any interference in nearby
		electronic equipment.
RF emissions	Class B	The models P01, P05 are suitable for used in domestic
CISPR11		establishment and in establishment directly connected
Harmonic emissions	Class A	to a low voltage power supply network which supplies
IEC 61000-3-2		buildings used for domestic purposes.
Voltage fluctuations	Complies	
/ flicker emissions		
IEC 61000-3-3		

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 model P08M 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
	test level	level	-guidance
Electrostatic	±8 kV contact	±8 kV contact	Floors should be wood, concrete or
discharge (ESD)	±2 kV, ±4 kV, ±8 kV,	±2 kV, ±4 kV, ±8 kV,	ceramic tile. If floors are covered with
IEC 61000-4-2	±15 kV air	±15 kV air	synthetic material, the relative
			humidity should be at least 30 %.
Electrical fast	±2kV for power	±2kV for power	Mains power quality should be that of a
transient/burst	supply lines	supply lines	typical commercial or hospital
IEC 61000-4-4	±1 kV for		environment.
	Input/output lines		
Voltage dips,	<5 % UT	<5 % UT	Mains power quality should be that of a
short	(>95% dip in UT.)	(>95% dip in UT.)	typical commercial or hospital
interruptions and	for 0.5 cycle	for 0.5 cycle	environment. If the user of the models
voltage	<5 % UT	<5 % UT	P01, P05 require continued operation
variations on	(>95% dip in UT)	(>95% dip in UT)	during power mains interruptions, it is
power supply	for 1 cycle	for 1 cycle	recommended that the models P01,
input	70% UT	70% UT	P05 powered from an uninterruptible
lines	(30% dip in UT)	(30% dip in UT)	power supply or a battery.
IEC 61000-4-11.	for 25/30 cycles	for 25/30 cycles	
	<5% UT	<5% UT	
	(>95 % dip in UT)	(>95 % dip in UT)	
	for 5/6 sec	for 5/6 sec	
Power frequency	30 A/m	30 A/m	Power frequency magnetic fields
(50/60 Hz)			should be at levels characteristic of a
magnetic field			typical location in a typical commercial
IEC 61000-4-8			or hospital environment.
NOTE UT is the a.c.	mains voltage prior to app	lication of the test level.	
Conducted RF IEC	3 Vrms	3 Vrms	Portable and mobile RF communications
61000-4-6	150 kHz to 80 MHz	150 kHz to 80 MHz	equipment should be used no closer to any
	6 Vrms in ISM and	6 Vrms in ISM and	part of the models P01, P05 including cables,
	amateur radio bands	amateur radio bands	than the recommended separation distance
			calculated from the equation applicable to the
			frequency of the transmitter.
			Recommended separation distance
			d=[3,5/V1]×P1/2
Radiated RF IEC	10 V/m	10 V/m	d=1.2×P1/2 80 MHz to 800 MHz
61000-4-3	80 MHz to 2.7 GHz.	80 MHz to 2.7 GHz	d=2.3×P1/2 800 MHz to 2.7 GHz
	385MHz- 5785MHz Test	385MHz- 5785MHz Test	where P is the maximum output power rating
	specifications for	specifications for	of the transmitter In watts (W) according to
	ENCLOSURE PORT	ENCLOSURE PORT	the transmitter manufacturer and d Is the

IMMUNITY to RF
wireless communication
equipment (Refer to
table 9 of IEC 60601-
1-2:2014)

IMMUNITY to RF wireless communication equipment (Refer to table 9 of IEC 60601-1-2:2014) recommended separation distance in meters (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:



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 model P08M is used exceeds the applicable RF compliance level above, the model P08M should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the model P08M.

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 models P08M

The model P08M are intended for use in electromagnetic environment in which radiated RF disturbances is controlled. The customer or the user of the model P08M can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the model P08M are 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			
	150kHz to 80MHz d=1.2×P1/2	80MHz to 800MHz d=1.2×P1/2	800MHz to 2,5GHz d=2.3×P1/2	
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) accordable 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.

CHANGE HISTORY		
Version	Description	Date
1.0	First release version	2022-03-29



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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.