

Hyper-Box Homecare

System

User Manual





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2 Preface

Application

This documentation applies to the product described as:

• Hyper-Box Homecare System

You will find this description on the nameplate on the rear of the device.

Firmware version

This documentation applies to the software version:

• SW Version 3.0 (or greater)

Subject to technical modification without prior warning.

Key to symbols used in this manual		
Keys, Hyper-Box labels and information in the display:	Keys, such as <i>Enter</i> , Hyper-Box Labels, such as <i>Air IN</i> , and information in the display, such as <i>Change Settings?</i> , are shown in bold, italic type.	

These products are covered by various USA and International Patents that can be found at: <u>www.aotinc.net/patents</u>



3 Intended Use

3.1 Intended Use

The **Hyper-Box** Homecare System is intended to be used for the treatment of acute and chronic wounds, such as:

- skin ulcerations due to diabetes, venous stasis, post surgical infections and gangrenous lesions
 decubitus ulcers
- decubitus ulcers
- amputations/infected stumps
- skin grafts
- o burns
- o frostbite

The device can be used in hospitals, nursing homes, clinics, as well as at the patient's home as an adjunct to standard wound therapy.

Warning!

Inadequate cleaning and disinfection of the *Hyper-Box Homecare Therapy* controller after use may lead to transmission of infectious disease.

Inadequate room ventilation, or Inappropriate venting of oxygen from the *Hyper-Box* Homecare System could lead to fire or explosion.

Never smoke or use a naked flame in the presence of oxygen.

To avoid risk of electric shock, this equipment must only be connected to a supply mains with protective earth.

Do not modify this equipment without authorization of the manufacturer.

Do not use in the presence of acute thrombophlebitis or DVT.



USA federal law restricts this device to sale or use by or on the order of a physician.

The *Hyper-Box* Homecare System should be operated by trained personnel under the direction of a qualified medical practitioner.



Technical Data 4 Configuration 4.1 4.1.1 Pneumatics Input flow > 10 L/min 5-60 Psi (0.3-4 Bar) Input Pressure 10 - 50 mbar Therapy Pressure 4.1.2 Special functions Automatic stand-by mode 4.1.3 Expected Service 10 Years l ife 4.1.4 Monitoring Operating time Operating pressure Therapy pressure too high Internal temperature too high Therapy complete 4.1.5 Physical data W x L x H of Single-Use 9 x 18 x 30 inches (23 Hyper-Box Chamber x 46 x 76 cm) W x L x H of Homecare 8 x 9 x 5 inches (20 Controller Unit x 23 x 13 cm) Total weight of Hyper-box 6.4 lbs (2.9 kg) Homecare Controller 4.1.6 Rating **Operational Temperature** 50 to 104°F (10 to 40°C) **Operational Humidity** < 80 % RH IP21 IP Rating (Protection against vertically falling water drops when controller is tilted up to 15°) Storage and transport 14 to 140°F (-10 to conditions 60°C) at 5 to 95 % RH 4.2 Power supply 100 - 230 VAC Input voltage to Power supply 50 - 60Hz Output voltage to Power 12 VDC supply < 10VA Power consumption



4.3 Compliance and approvals

C€₀₀₅₀

The *Hyper-Box* Homecare System complies with the requirements of directive 93/42/EEC concerning Medical Devices and therefore bears the CE mark.

The *Hyper-Box Homecare System* is classified as protection class II, Type BF, steady state.

The *Hyper-Box* Homecare System complies with the following International standards:

IEC 60601-1 - Electrical Safety IEC 60601-1-2 - Electromagnetic Compatibility

4.4 Manufacturer's declaration

Guidance and manufacturer's declaration – electromagnetic emissions

The *Hyper-Box Homecare System* is intended for use in the electromagnetic environment specified below. The customer or the user of the Hyper-Box Homecare controller should assure that it is used in such an environment.

Emissions test	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11	Group 1	The Hyper-Box Homecare Therapy controller 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 Hyper-Box Homecare Therapy controller is suitable for use in all establishments, including
Harmonic emissions IEC 61000-3-2	Class A	domestic establishments and those directly connected to the public low-voltage power supply
Voltage fluctuations/flicker emissions IEC 61000-3-3	Complies	network that supplies buildings used for domestic purposes.

Immunity tests	EN 60601 Test level	Compliance level	Electromagnetic environment - guidance
Electrostatic discharge (ESD)	+/- 8 kV contact +/- 15 kV air	+/- 8 kV contact +/- 15 kV air	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30 %.
IEC 61000-4-2			
Electrical fast transient/burst	+/- 2 kV for power supply lines	+/- 2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
IEC 61000-4-4	+/- 1 kV for input/output lines	+/- 1 kV for input/output lines	
Surge	+/- 1 kV differential mode	+/- 1 kV differential mode	Mains power quality should be that of a typical commercial or hospital environment.
IEC 61000-4-5	+/- 2 kV common mode	+/- 2 kV common mode	
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	<pre>< 5 % UT (> 95 % dip in UT) for 0.5 cycle 40 % UT (60 % dip in UT) for 5 cycles 70 % UT (30 % dip in UT) for 25 cycles < 5 % UT (> 95 % dip in UT) for 5 secs.</pre>	< 5 % U _T (> 95 % dip in U _T) for 0.5 cycle 40 % U _T (60 % dip in U _T) for 5 cycles 70 % U _T (30 % dip in U _T) for 25 cycles < 5 % U _T (> 95 % dip in U _T) for 5 secs.	Mains power quality should be that of a typical commercial or hospital environment. If the user of the Hyper-Box Homecare controller requires continued operation during power mains interruptions, it is recommended that the Hyper-Box with O2 controller be powered from an uninterruptible power supply or a battery.
Power frequency (50/60Hz) magnetic field IEC 61000-4-8	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment

mmunity tests	EN 60601 Test level	Compliance level	Electromagnetic environment - guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the Hyper-Box Homecare controller, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
		0.) (Recommended separation distance:
Conducted RF EC 61000-4-6	3 V rms 150 kHz to 80 MHz	3 V rms	d = 1.2 x root P
Radiated RF	3 V/m	3 V/m	d = 1.2 x root P 80 MHz to 800 MHz
EC 61000-4-3	80 MHz to 2.5 GHz		d = 2.3 x root P 800 MHz to 2.5 GHz
			Where <i>P</i> is the maximum output power rating in 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.
NOTE 1: At 80 MHz a	and 800 MHz, the higher freq	uency range applies.	
NOTE 2: These guide structures, objects ar		uations. Electromagnetic	propagation is affected by absorption and reflection from
amateur ra electromag measured compliance	dio, AM and FM radio broadd gnetic environment due to fixe field strength in the location i e level above, the Hyper-Box	cast and TV broadcast ca ed RF transmitters, and el n which the Hyper-Box Ho Homecare controller sho	radio (cellular/cordless) telephones and land mobile radios nnot be predicted theoretically with accuracy. To access the ectromagnetic site survey should be considered. If the omecare controller is used exceeds the applicable RF uld b observed to verify normal operation. If abnormal /, such as reorienting or relocating the Hyper-Box Homecar

P/N: G00202

Therapy controller					
			ment in which radiated RF disturbance		
			elp prevent electromagnet interference		
			nt (transmitters) and the <i>Hyper-Box</i>		
Homecare Therapy controller	as recommended below, accord	ding to the maximum output power	of the communications equipment.		
Separation distance according to frequency of transmitter					
Rated maximum output	m				
power of transmitter (P)	150 kHz to 80 MHz	80 MHz to 800 MHz	800 MHz to 2.5 GHz		
W	d= 1.2 x root P	d= 1.2 x root P	2.3 x root P		
0.01	0.1	0.1	0.2		
0.1	0.4	0.4	0.7		
1	1.2	1.2	2.3		
10	3.8	3.8	7.3		
100	12	12	23		

calculated using the equation applicable to the frequency of the transmitter. *P* is the maximum output power rating of the transmitter in watts (W) as declared by the transmitter manufacturer.

NOTE 1: At 80 MHz and 800 MHz, the separation distance fort he 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.



4.5	Device labels and symbols		labels and symbols can be found on x Homecare System:
		Ò	<i>Power</i> switch: On/Off
			Navigation and value setting button <i>Up</i>
			Navigation and value setting button Down
		C	<i>Enter</i> button: change menu level, apply changed value and acknowledge of alarms
			DC Power Supply input: 12 VDC/800mA
		SN	Serial number
		\triangle	Caution
			No smoking!
			Do not use oil or grease!
		(Single Use - Do Not Reuse
		C€ 0050	CE number
		★	Applied part type BF, according to EN 60601-1

Topical Wound



Double Insulated



CSA & UL Approval

Recycle: Electronic Equipment



Consult instructions for use



Manufacturer



Humidity Limitation



Temperature Limitation



Do Not Use if Package is Damaged



Keep dry



Fragile; Handle with care

This Way Up

5 Safety Instructions

5.1 Symbol for warnings cautions and notes This triangular hazard symbol is used to draw your attention specifically to the remaining dangers associated with proper use and to emphasize important technical requirements.



Warnings, Cautions or Instructions, to prevent potential injury or damage.

5.2 Personnel

Caution

The *Hyper-Box* Homecare System should only be operated by personnel with suitable training under the order of a physician.

5.3 Basic safety information and warnings

(I) Warning!

The *Hyper-Box* Homecare System should only be used with medical grade oxygen from a Cylinder, Piping system, Liquid Oxygen System or Concentrator.

If utilizing a Liquid Oxygen System (LOX) with the device, follow all rules pertaining to the safe handling of liquid oxygen, as detailed in Appendix B

Only use clean, oil-free DISS hoses and connections to connect the **Oxygen Source** to the Hyper-Box Homecare controller.

Inadequate room ventilation, or inappropriate venting of oxygen from the *Hyper-Box* Homecare System could lead to fire or explosion.

Never smoke or use an open flame in the presence of oxygen.

To disinfect the *Hyper-Box* Homecare Controller - we recommend use of the specified cleaners detailed in section 9 - *Care and Maintenance*, of this manual.

If the *Hyper-Box Homecare System* is damaged in any way, or the supply of oxygen can no longer be guaranteed - stop treatment with the device immediately.

5.4 Potential Potential risks associated with the use of this device may include:

- infection, if proper cleaning is not performed
- burn, if caution is not taken with the liquid oxygen or electrical fittings
- tissue damage, if instructions for use are not followed

6 Preparing for Use

6.1 Personnel

6.2 Hyper-Box Homecare Therapy Single-Use Chamber **Caution** The *Hyper-Box* Homecare System should only be operated by personnel with suitable training under the order of a physician.

\Lambda Warning!

The chamber is for single use only. **Do not** reuse, reprocess or disinfect. Reuse, reprocessing or disinfection may compromise the structural integrity of the device and/or lead to device failure which, in turn, may result in patient harm or illness. Reuse, reprocessing or disinfection may also create a risk of contamination of the device and/or cause patient infection or cross-infection, including, but not limited to, the transmission of infectious disease(s) from one patient to another. Contamination of the device may lead to harm or illness to the patient.

After use, dispose of product and packaging in accordance with administrative and/or local government policy.



Powering off the unit by pressing the On/Off button or removing the Quick Connect hose, will relieve cuff pressure if required to remove patient's limb from the Single Use Chamber at any time during Therapy.



6.3 Items supplied with the Hyper-Box Homecare Therapy Single-Use Chamber and Hyper-Box Homecare System Single Use Chamber and Controller





DISS Oxygen line



Power supply



Aeroneb Solo Humidifier and Cable



Interface Quick Connect Hose



User manual

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6.4 Set up and assembly

The set up and assembly is detailed below.

- 6.5 **Pre-Cleaning** Prior to use the external surfaces of the *Hyper-Box Homecare Therapy* controller and accessories should be cleaned and disinfected with an effective intermediate level disinfectant as referenced is section 9.
- 6.6 Connecting the Power supply

Take the Power supply, Select and affix the appropriate power plug provided. Connect the 12 DC output to the DC socket on the rear of the **Hyper-Box** Homecare Therapy controller.

Ensure to attach the DC Power cable to the P-Clip.

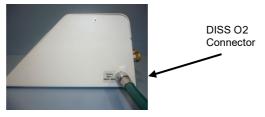
DC Power supply cable and input connector



- 6.6.1 Power Supply Input voltage
- 6.7 Connecting the DISS Oxygen hose to the controller unit

The power supply should be connected to any 100 -240 VAC 50/60 Hz outlet.

Connect the DISS oxygen hose to the DISS connector located on the Side of the controller unit



Warning!

The *Hyper-Box Homecare System* should only be used with medical grade oxygen from a Cylinder, Piping system, Liquid Oxygen System or Concentrator

Do not obstruct the vent port on the *Hyper-Box Homecare System* homecare controller.

Use only in a well-ventilated room. Inadequate room ventilation, or inappropriate venting of oxygen from the *Hyper-Box Homecare System* could lead to fire or explosion.

Never smoke or use an open flame in the presence of oxygen.

If utilizing a Liquid Oxygen System (LOX) with the device, follow all rules pertaining to the safe handling of liquid oxygen, as detailed in Appendix B

Only use clean, oil-free DISS hoses and connections to connect the Oxygen Source to the **Hyper-Box** Homecare Therapy controller.

Installation instructions:

- Connect the alternate end of the DISS oxygen hose to the regulator/flowmeter outlet on the oxygen cylinder or LOX system.
- Slowly open the main valve on the oxygen cylinder or LOX system.
- 3. Set the flowmeter to 10 lpm

6.9 Connecting the DISS Oxygen hose to the hospital's piped medical gas supply or Concentrator

Connecting the

Oxygen (LOX)

system, High Pressure Cylinder

to a Liquid

DISS Oxygen hose

6.8

Installation instructions:

- Connect the DISS oxygen hose to the DISS oxygen outlet of the flowmeter connected to the piped wall supply or Concentrator.
- 2. Set the flowmeter to 10 lpm.



6.10 Setting up the AeroNeb Solo Humidifier

Take the AeroNeb humidifier and connect the interface cable. Then connect the interface cable to the humidifier output on the rear of the controller.



Unpack the Single-Use chamber assembly and install the humidifier into the Single-Use chamber as shown below:



It is recommended to keep humidifier facing upright during treatments.

Fill the AeroNeb cup with sterile demineralized (distilled) water.



The Use of fluids other than sterile distilled water for humidification may cause damage to the humidifier and *Hyper-Box* Homecare System

After each treatment, dispose of the unused water utilizing standard procedures

The AeroNeb Humidifier is Single Patient Use and should be disposed of between patients.



6.11 Connecting the Single-Use chamber to the controller

Take the Interface Hose and connect the Quick connect plug to the Quick connect socket on the Controller unit (the smaller plug). Connect the Other end of the Hose to the Quick connect socket on the Single-use chamber as shown (larger plug)

The Quick connect plug and sockets are keyed to guarantee correct orientation and connection.





7 Applying the Therapy

7.1	Basic description	The <i>Hyper-Box Homecare System</i> is intended as adjunctive therapy to standard wound care for acute or chronic wounds.
		The <i>Hyper-Box Homecare Therapy</i> controller regulates the humidified oxygen pressure and the duration of treatment inside the Single-Use extremity chamber in accordance with the physician's orders.
		The Single-Use chamber should be disposed of between treatments.
7.2	Preparing the wound	It is recommended to remove all bandages, dressings, creams and ointments from the wound prior to treatment.
		Wounds should be cleaned, and debrided if required, per standard wound care protocols, prior to commencing treatment with this device
7.3	Recommended Treatment duration	Recommended treatment durations are:
		• 60 – 90 mins. once a day, five days per week

7.4 Infected wounds It is recommended that all infected wounds (including those infected with antibiotic-resistant strains, such as MRSA) be treated utilizing the institutions' standard operating procedures in conjunction with the therapy.



7.5 Applying the therapy utilizing the Single Use Chamber

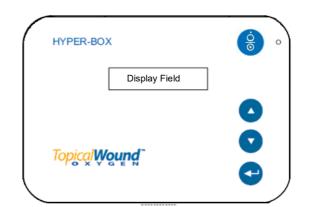
(2)

- 1. Set the desired therapy pressure and duration as prescribed by the physician as detailed in section 8 of this manual.
- 2. Press the "Power On" button for 2 seconds
- Select "Inflate Bag" by pressing the enter key on the controller unit. The deflated chamber will slowly inflate and become ridged. "Inflating Bag" will be displayed during this process.
- 4. Insert the patient's limb through the patient inlet, making sure that the chamber inner sides do not come in contact with the wound.
- Select "Inflate Cuff" by pressing the enter key on the controller unit. The deflated patient cuff will inflate around the patient's limb and create a seal. "Inflating Cuff" will be displayed during this process.
- Select "Start Therapy" by pressing the enter key on the controller unit. The therapy will now be applied to the patient per the physician's orders.
- The therapy can be paused if required by pressing the enter key during the pressurizing cycle of the therapy. "Therapy Paused Up to continue" will be displayed. To resume the treatment press the "Up" button to continue.
- 8. The therapy will automatically stop once the set treatment duration has completed and the Patient Cuff will deflate.
- 9. Remove the patient's limb from the chamber.
- 10. Dispose of the single use chamber.
- **7.6 Aftercare** After treatment, the wound should be dressed utilizing the desired dressings as recommended by the clinician and following standard wound care procedures.



8 Controller Unit Operation

8.1 Description of operating controls



Display Field	The display field provides information on the monitoring values for the <i>Hyper-</i> <i>Box Homecare System</i> and is used to enter setting values
LED	The LED provides a visual alarm
On/Off	Use the On/Off switch to switch the device on, or power down to the standby mode.
Navigation button <i>UP</i>	Use the navigation button Up to scroll up a menu
Navigation button DOWN	Use the navigation button Down to scroll down a menu
Enter	The <i>Enter</i> button is used to move between menu levels and to acknowledge alarms.



8.2	Personnel	
		The Hyper-Box Homecare System should only be operated by personnel with suitable training under the order of a physician.
8.3	Switching the device on/off	Check that all cables and tubes are correctly installed. Start the device by pressing the On/Off button
8.4	The start screen	When the <i>Hyper-Box</i> is switched on using the <i>On/Off</i> button, the welcome screen appears. After two seconds the message "Inflate Bag" is displayed.
		The buzzer will bleep upon switch on.
8.5	Operating	The message "Inflate Bag" is displayed.
	Screens	Press the <i>Enter</i> key to inflate the chamber.
		The message "Inflating Bag" is displayed.
		The message "Inflate Cuff" is then displayed.
		The message "Inflate Cuff" is then displayed. Press the <i>Enter</i> key to inflate the patient cuff.
		Press the <i>Enter</i> key to inflate the patient cuff.
		Press the <i>Enter</i> key to inflate the patient cuff. The message " Inflating Cuff " is displayed.
		Press the <i>Enter</i> key to inflate the patient cuff. The message "Inflating Cuff" is displayed. The message "Start Therapy?" is then displayed
		Press the <i>Enter</i> key to inflate the patient cuff. The message "Inflating Cuff" is displayed. The message "Start Therapy?" is then displayed Press <i>Enter</i> to commence the therapy. During the therapy the cyclical therapy pressure and a
8.6	Pausing Therapy	Press the <i>Enter</i> key to inflate the patient cuff. The message "Inflating Cuff" is displayed. The message "Start Therapy?" is then displayed Press <i>Enter</i> to commence the therapy. During the therapy the cyclical therapy pressure and a countdown of therapy time is displayed' Pressing the <i>On/Off</i> key will cause the therapy to stop
8.6	Pausing Therapy	Press the <i>Enter</i> key to inflate the patient cuff. The message "Inflating Cuff" is displayed. The message "Start Therapy?" is then displayed Press <i>Enter</i> to commence the therapy. During the therapy the cyclical therapy pressure and a countdown of therapy time is displayed' Pressing the <i>On/Off</i> key will cause the therapy to stop and patient cuff to deflate. To pause the treatment press the <i>Enter</i> key during the
8.6	Pausing Therapy Therapy Complete screen	 Press the <i>Enter</i> key to inflate the patient cuff. The message "Inflating Cuff" is displayed. The message "Start Therapy?" is then displayed Press <i>Enter</i> to commence the therapy. During the therapy the cyclical therapy pressure and a countdown of therapy time is displayed' Pressing the <i>On/Off</i> key will cause the therapy to stop and patient cuff to deflate. To pause the treatment press the <i>Enter</i> key during the pressuring cycle of the therapy.



8.8	The monitoring and settings menu		monitoring and settings menu displays a number of ctable or monitored parameters. These include:
		1.	The therapy pressure in mbar
		2.	The therapy duration in minutes
		3.	The accumulative therapy hours
			can scroll through these menus, or select another lay value, by pressing the navigation buttons Up and vn .
8.9	Changing Settings for Therapy		can set the therapy pressure and duration as scribed by the physician as follows;
	Pressure and Duration, and for viewing Running hours	weld	t the device by pressing the On/Off button, The come screen appears. After two seconds the Inflate message is displayed.
			ss the <i>Up</i> and <i>Down</i> arrow together to progress into the ings and calibration menu.
			the <i>Up</i> and <i>Down</i> arrow to adjust prescribed therapy ssure and then press <i>Enter</i> .
			the <i>Up</i> and <i>Down</i> arrow to adjust prescribed therapy ation and then press <i>Enter</i> .
			se values are now stored in memory and will be ined until changed using this process.
		The	accumulative therapy running hours are now displayed
		You	can now exit by pressing the On/Off button,
			ou wish to zero the pressure transducers, press the <i>Up</i> w instead of the <i>On/Off</i> button,
		and	te sure the Quick connect tubing is disconnected press <i>Enter</i> . The pressure transducers are now ped.
		You	should now exit by Pressing the On/Off button,
8.10	The alarm screen		alarm is triggered, a corresponding message will ear in the display.
			ns are divided into two priority classes:

- High priority alarm
- Medium priority alarm



8.10.1 Alarm Messages	An alarm is displayed when a certain alarm condition is met. The alarm display ends as soon as the alarm condition no longer exists.	
'Check for leaks' Alarm	If the set pressure in the system is not reached within 3 minutes ' Check for leaks ' message will be displayed.	
	Che	ck the following:
	1.	Oxygen hose is connected and gas supply is switched on.
	2.	The interface hose is connected to controller and single use chamber.
	3.	The cuff is inflated and there is no leak around the limb.
	4.	The humidifier is inserted into the single use chamber.
	5.	Replace the single use chamber.
		e the issue is resolved restarting the unit will resume nal operation.
Loss of Pressure		stem pressure drops significantly during use the unit will matically shut off.
	Che	ck the following:
	1.	The gas supply is switched off.
	2.	The interface hose has not got disconnected connected from the controller or single use chamber.
	3.	The humidifier has come out or not inserted fully.
	4.	The limb was removed from the single use chamber during therapy.
	5.	The single use chamber has a leak. Replace the single use chamber.
		e the issue is resolved restarting the unit will resume nal operation.
8.10.2 Warnings		Warning!

Ignoring any alarm may interfere with the functionality of the device and could result in patient injury.

9 Care and Cleaning

9.1 Personnel

Caution

The *Hyper-Box* Homecare System should only be operated and maintained by personnel with suitable training on the order of a physician.

9.2 Guidelines for care and maintenance To ensure the safe and effective functionality of the *Hyper-Box Homecare System* controller it is important that the device is cleaned and maintained according to these instructions. Only use spare parts and accessories recommended by the manufacturer.

Caution

Guidelines and maintenance instructions provided by the manufacturer should be followed.

Inadequate cleaning and disinfection of the *Hyper-Box Homecare Therapy* controller and its accessories after use may lead to transmission of infectious disease.

Always observe and follow the manufacturer's instructions for use for the cleaning/disinfectant substance.

9.3 Limitations on Reprocessing: Due to the product design and the materials used, no definite limit to the maximum number of performable Cleaning and Disinfection processing cycles can be specified. The service life of the medical device is determined by their function and careful handling.



9.4 Cleaning and Disinfection

Dispose of the Single-Use chamber after use.

After each treatment, dispose of the unused water utilizing standard procedures

The AeroNeb Humidifier is Single Patient Use and should be disposed of between patients.

The external surfaces of the *Hyper-Box* Homecare Therapy controller and accessories should be cleaned and disinfected after every use with an effective intermediate level disinfectant.

Even antibiotic-resistant organisms, such as MRSA, are effectively destroyed by the intermediate level disinfectants listed below.

Clean and Disinfect all external surfaces of the Controller, including gas connections and hose, Quick Connect Interface hose, Humidifier cable and plug, Power supply and power cord.

The following intermediate level cleaner/disinfectants (or equivalents) are recommended for this purpose:

Alpha Guard GF

Dr. Deppe Laborities GmbH, Germany

Bacillol 30 Tissue Wipes

BODE Chemie GmbH, Germany

Professional AMPHYL® II Disinfectant Deodorant Spray

Reckitt Benckiser Inc., USA.

Cleaning and Disinfection Instructions

 9.5 Transport No particular requirements. It is recommended that instruments are reprocessed as soon as is reasonably practical following use.
 9.6 Preparation for Cleaning: Disconnect reusable chamber from the Quick Connect Hose and the Humidifier and dispose of Chamber. Disconnect Quick connect Hose from Controller unit. Disconnect the Humidifier cable from the controller. Unplug the power supply unit.



 9.7 Disinfection with Amphyl II Deodorant Spray
 Professional AMPHYL® II Disinfectant Deodorant Spray kills microorganisms on environmental surfaces. To Disinfect: Hold can upright 6" to 8" from surface. Spray precleaned surfaces 2 to 3 seconds until covered with mist. Spray Controller unit, Quick connect hose, power supply and cord, Humidifier cable and Oxygen hose.

Allow to stand for 10 minutes to air dry.

 9.8
 Disinfection with AlphGuard Solution:
 Disinfectant (AlphGuard Solution) must be used in accordance with the manufacturer's labeled instructions. Gloves should be worn.

Application Procedure:

1. Select appropriate concentration per the table below, such as 1.5% (15 mL AlphaGuard / 1L water).

Concentration	Exposure Time (Minutes)
1.50%	15
1.00%	30
0.50%	60
0.25%	240

2. Mix AlphaGuard with water in a bottle per the concentration ratio desired.

3. Use a cloth or paper towel to apply solution to all external surfaces of the Controller unit. Also apply solution to the external surfaces of the Quick connect hose, power supply and cord, humidifier cable and Oxygen hose.

4. Observe required exposure time to achieve disinfection.



9.9	Disinfection with Bacillol 30 Tissue Wipes:	Disinfectant (Bacillol 30 Tissue Wipes) must be used in accordance with the manufacturer's labeled instructions. Gloves should be worn.
		Application Procedure:
		1. Thoroughly wipe all external surfaces of the Controller unit, including; the Quick connect hose, power supply and cord, Humidifier cable, Gas connections and Oxygen hose with the Bacillol 30 Tissues.
		Make sure to completely wet all surfaces to achieve best disinfection results.
		3. Dispose of used tissues after use.
		4. Allow for 30 min exposure time to guarantee disinfection.
9.10	Inspection and Functional Testing:	Optically inspect for cleanliness. Repeat cleaning and Disinfection procedure if necessary. Prior to use perform functional testing and setup according to Section 6 of the User Manual.
9.11	Packaging:	A standard packaging material may be used.
9.12	Storage:	Store disinfected device in a dry, clean, and dust-free environment at moderate temperatures of between 5°C to 40°C.



Preventive Maintenance 10

10.1 Scheduled Preventive Maintenance

Refer to Hyper-box Service Manual (P/N G00316) for further information on Preventative Maintenance and Technical Service instructions.

Interval	Part	Procedure
After Each Patient Therapy	Single-Use Chamber	After each therapy dispose of the Single- Use chamber following standard local procedures. The chamber is for single use only. Do not reuse, reprocess or disinfect.
After Each Patient Treatment	Humidifier	After each treatment, dispose of the unused water utilizing standard procedures.
		The AeroNeb Humidifier is Single Patient Use and should be disposed of between patients.
<u> </u>	1	
Annually	Exhaust Muffler	Replace the chamber vent muffler (G00141)

 Manufacturer's address	AOTI Ltd., Unit 20, Glenrock Business Park, Ballybane, Galway. Ireland.
 Technical support & customer service	AOTI Ltd., Unit 20, Glenrock Business Park, Ballybane, Galway. Ireland. customerservice@aotinc.net
 Address for orders	AOTI Ltd., Unit 20, Glenrock Business Park, Ballybane, Galway. Ireland. customerservice@aotinc.net Tel.: +353 9166 0310

11 Accessories and Spare Parts

1.1 Accessories and	Item	Order number
spare parts	Hyper-Box Homecare System	G00001-2
	Alpha Guard disinfectant/cleaner	G00005
	Single-use HyperBox Chambers (Case of 20) Medium	G00020
	Single-use HyperBox Chambers (Case of 20) Large	G00061
	Humidifier	G00113
	Humidifier Cable	G00114
	Quick Connect Interface Hose	G00126
	Muffler Sintered Bronze, 1/4",	G00141
	Oxygen Supply Hose ISO, DISS Nut - DISS Handtight (White)	G00191
	Oxygen Supply Hose US, DISS Nut - DISS Handtight (Green).	G00192
	Power Supply, 100 - 240 VAC, O/P 12v, Euro	G00250
	Power Supply, 100 - 240 VAC, O/P 12v, US/Japan	G00251
	Power Supply, 100 - 240 VAC, O/P 12v, UK	G00252
	Service Manual	G00316
	Professional AMPHYL® II Disinfectant Deodorant Spray, Box of 12.	G00335
	Bacillol 30 Tissue Wipes	G00336



12 Disposal

The operator is responsible for the disposal of the device. The operator must either ...

- Deliver the device, free of charge and duty paid, to the manufacturer for disposal or
- Surrender the device to a licensed private or public collection company or
- Competently recycle the parts/dispose of them properly.

Where an operator chooses to dispose of the device himself, the disposal regulations are specified for each country and laid down in the appropriate laws and regulations. Please consult the responsible authorities for instructions on what is required.

With this in mind, all waste is to be recycled or disposed of:

- Without any risk to human health
- Without employing procedures or methods which cause damage to the environment in particular water, air, earth, flora and fauna
- Without causing any noise or odour nuisance
- Without detriment to the surroundings or landscape.



13 Ap	opendix A: Abbreviations and Glossary
Α	
A	Ampere
AC	Alternating Current
AT	Ampere Slow Blow (fuse)
B	
Bar	1 bar = 14.50 psi
<u>°C</u>	Dumun Ouluin
-0	Degrees Celsius Converting Celsius (C) to Fahrenheit (F):
_	F = 9*C/5 + 32
D dBA	Desikels recoursed with a filter
DC	Decibels measured with a filter Direct Current
 °F	Degrees Fahrenheit
	Converting Fahrenheit (F) to Celsius (C):
	C = (F-32)*5/9
н	
Hz	Hertz (1 Hz = 1 s ⁻¹)
1	
IP	Safety class according to ISO standards
_L	
L	Litres
Lbs	Pounds Litres per minute of flow
Lpm	Littles per minute of now
Μ	
Max., ma	
Min	Minute
min.	Minimum Millimatros of moroury processo
mmHG -	Millimetres of mercury pressure
P psi	Pressure pounds per square inch (1 bar = 14.50 psi)
psi	Pressure pounds per square mon (1 bar – 14.50 psi)
R	
RH	Relative humidity
<u>T</u> TF	
TF	Technical fault
V	
V	Volt
VA	Apparent power of device
VAC	Volt Alternating Current
VDC	Volt Direct Current

14 Appendix B: LOX Use

PATIENT USER TRAINING CARD

USE OF MEDICAL LIQUID OXYGEN VESSELS IN THE HOME

