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The Next Step in Patient Transfer



AEROPOD[®]
User Manual



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Contents

Company Introduction	3
Introduction	4
Intended Use Statement	6
Aviation Compliance	7
Key Features	8
 Product Features	 9
• Safety Harness	10
• Hinge System	10
• Structure	10
• Air Ventilation and Filter System	11
• Access Ports	11
• Connector Ports	11
 Instructions on use.....	 12
• Access Ports	12
• Changing Gloves.....	12
• Using supply bags.....	13
• Using connector ports.....	14
 Ventilation System	 15
• Negative Pressure	16
• Positive Pressure	17
 Assembly Instructions.....	 18
• Patient Loading	19
• Patient Unloading	20
 Decontamination Protocol	 21
Emergency Procedures	23
User Requirements	24
Organisational Requirements	24
Product Specification	25
Storage	27
Assembly and Patient Loading Checklist ...	28
Decontamination Record	29



Company Introduction

BBL Protect was established in 2020. We worked through the pandemic learning from the practice development of professionals and organisations in the fight against the Covid-19 virus including working within organisations on infection prevention and control standards and the needs of healthcare professionals.

The aim was to create a range of devices and products that meet the challenges of all situations providing protection for the environment, attending staff and others from cross-contamination.

The devices are designed for use within healthcare settings for the internal movement of patients and also for the emergency and pre-planned movement of patients to healthcare settings.



Introduction

AeroPod is a Personal Isolation Device designed for the movement of persons with infectious diseases, contaminants or that need protecting from the external environment around them. AeroPod has been designed for use on any kind of stretcher and for use on rotary and fixed wing aircraft, road ambulances and other vehicles.

AeroPod is registered as a class 1 medical device under the European MDR Guidelines and has full aviation compliance for use on rotary, CS23 and CS25 aircraft.

The Pod is designed for use with an external airflow supply operating with a minimum flow of 15 air exchanges per hour. Given the volume of the Pod, this generates a flow rate of 165 litres per hour. The unit can also be used with negative pressure drawing air out of the unit through the airflow system, and air in through the passive air filter system.

The Pod has been designed to minimise weight whilst maintaining the highest level of efficacy possible.

AeroPod has a simple to use harness system which allows securing of the patient directly to the stretcher, ensuring maximum safety.

This user manual will provide all necessary information for the use of the AeroPod.

The information in this manual is for reference purposes only and for the use of AeroPod only.

This information should be used in line with organisational guidelines, policies and procedures.



AeroPod



Important



- **DO NOT LEAVE PATIENTS UNATTENDED WHILST IN AEROPOD**
- **DO NOT drop the device onto surfaces or ground**
- **DO NOT leave device contaminated after use**
- **DO NOT leave waste items in the device after use**
- **DO NOT leave used items attached to the device after use**
- **DO NOT leave device unsecured to the stretcher**
- **DO NOT undo structural supports without first fully undoing the zip**
- **DO NOT use the device to carry paperwork required during transport**
- **DO NOT balance anything on top of the device**

Symbol Meanings



Signifies European technical conformity



Includes the address of the device manufacturer



Indicates the date when the device was manufactured



Indicates the manufacturers batch number for identification



Consult instructions for use



Indicates the Unique Device Identification



Indicates the item is a medical device

Intended Use Statement

AeroPod is a Personal Isolation Device for the transportation of persons with infectious diseases, chemical contamination or who are immunocompromised, from the first point of contact to required treatment centre or for strategic movement of patients between treatment centres.



Aviation Compliance

AeroPod meets the specifications as set out in CS25.853 including the Flammability Regulations (FR) for materials.

Under Aviation regulations a minor modification or STC approval will be required for the Pod to be used on each stretcher type and aircraft combination. The type of approval required will depend on the aircraft type.

Use on Pressurised Aircraft

When used on pressurised aircraft the patient oxygen mask must be connected to the in-line oxygen filter which in turn must be connected to an external oxygen cylinder for it to be available to the patient in the event of sudden aircraft decompression.

In the event of a sudden aircraft decompression the patient will need to put on the oxygen mask and the attending staff must switch on the oxygen cylinder allowing the patient to draw on oxygen.

Where the patient is not able to do this themselves the attending staff must fit the patients mask for them using the available glove ports prior to switching on the oxygen feed.



Key Features

Medical Compliance

AeroPod is registered as Class 1 Medical Device and holds the relevant CE mark in Europe.

Compatibility

AeroPod has a simple system for safely securing the patient directly to the stretcher on ambulances, aircraft and helicopters.

In Transport Care

AeroPod has 8 connector ports for IV lines, monitoring cables and other equipment. A filtered in-line oxygen connector is available for direct oxygen feed to the patient.

Quick Deployment

AeroPod is intuitive in its deployment and is designed to be quickly assembled for use.

Battery Powered Airflow

The air purifying device allows for up to 10 hours use. The battery can also be changed whilst the Pod is in use.

Aviation Compliance

AeroPod has full aviation compliance under EASA, CAA and FDA regulations for use on aircraft.

Patient Care

Access to the patient is through 8 capped ports. Glove rings allow for the use of disposable gloves and can be used with bags for supply of equipment into the sealed pod.

Dual Protection System

AeroPod is driven by an external air purifying respirator system offering both positive and negative pressure with the added benefit of attaching to the stretcher or worn round the waist by staff.

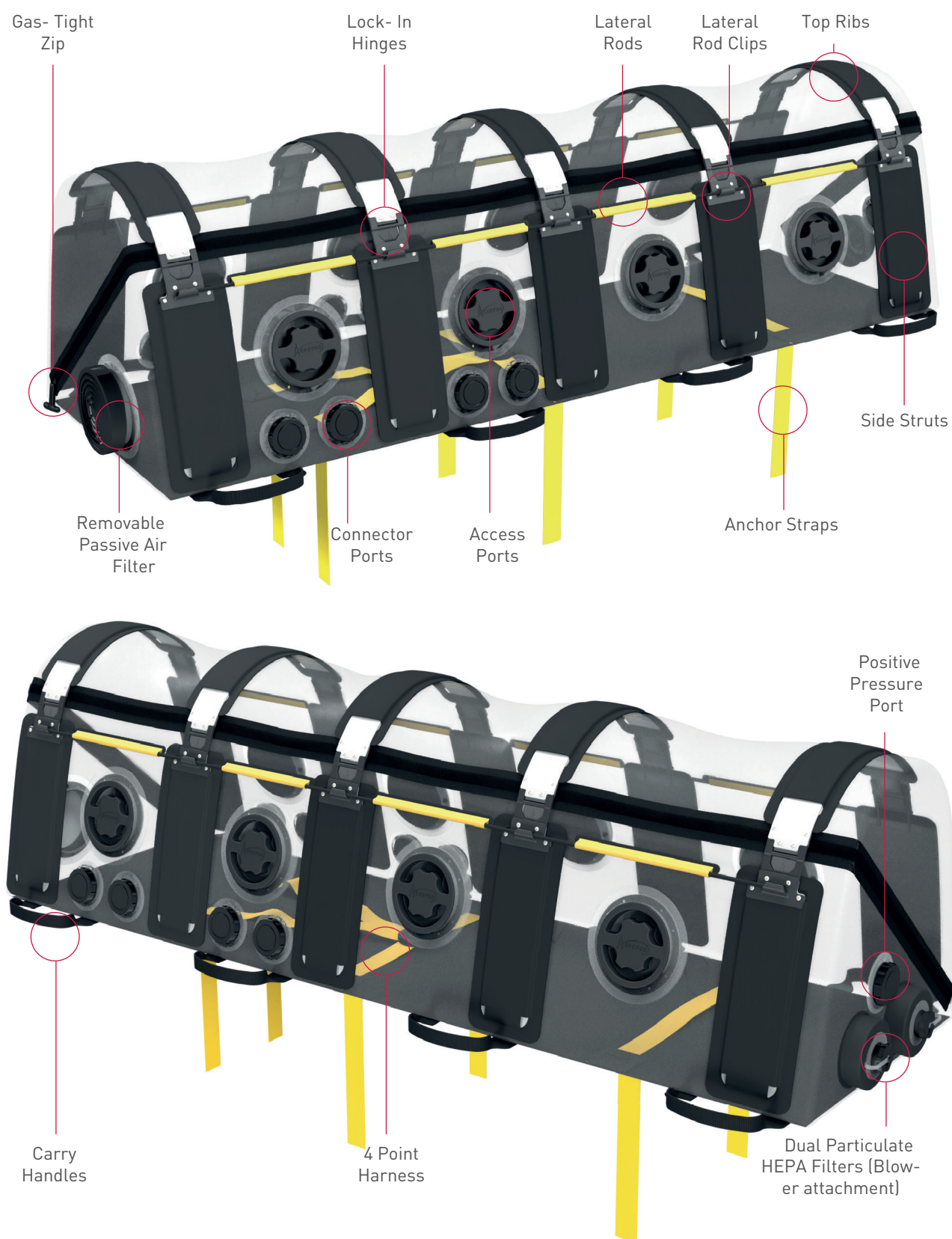
Reusable

The unit is simple and easy to clean for re-use multiple times. The only single use items are the filters, the gloves in the access ports, the supply bags and the cones in the connector ports.

Recyclable

The AeroPod is 90% recyclable depending on prior use and decontamination processes.

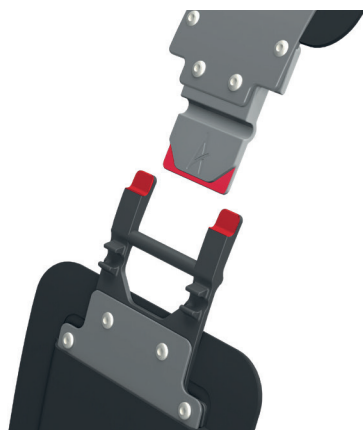
Product Features



Safety Harness

An aviation standard safety harness secures the patient directly to any stretcher.

Straps looped around the stretcher and using the double- D rings secure the pod to the stretcher.



Hinge System

The hinge system is designed to offer a quick and secure method of opening and locking the AeroPod.

Once zipped up the hinges cannot come undone, providing a safe and secure structure.

Structure

The AeroPod product structure is manufactured from highly engineered plastics which are highly resistant to heat, chemicals and are scratch resistant.

Operating temperature of the material is -60 to 175 degrees Celsius.



Air Ventilation and Filter System

The device offers both positive and negative pressure using an APR HEPA system filtering the air into and out of the device providing more than 15 complete air exchanges per hour. The battery offers up to 10 hours of uninterrupted power.



Access Ports

8 ports offer access to the patient during transport via a gloves system. A bag system can also be connected to the glove rings to supply items into the AeroPod.

Ports must remain sealed at all times during transfers, when being accessed.

Connector Ports

The connector ports allow for the pass through of IV lines and electrical monitoring equipment via conical sheaths which are taped to create a seal.

AeroPod uses a simple but recognisable system of a PVC cone secured in place by a compression screw cap.

Tubes and wires can be passed through this once the end of the cone has been cut off to the appropriate size and secured around the tube with strong tape to seal.

The oxygen ports have an in-line filter preventing any back flow contagion from inside the Pod. This connects directly to an oxygen tank outside the Pod and via a tube and mask to the patient.



Instructions on Use

Access Ports

Ports must remain sealed at all times during transfers, when being accessed.

The access ports have four key parts:

Housing - is glass filled nylon coated in flexible TPU allowing them to be welded to the shell of the device ensuring a secure seal.

External Cap - This is operated from outside the device and is opened and closed through a quarter turn.

Internal Cap - The internal cap can be operated from outside the unit through a quarter turn.

The external and internal caps are both secured through a bayonet fitting and are released via a quarter turn to the left and secured through replacing and rotating a quarter turn to the right.

Glove ring - The glove ring attaches into the housing once the external cap has been removed. The gloves are stretched onto the ring and secured by a heavy duty band.

Once the glove ring is secured in place with a quarter turn the internal cap can then be removed and the glove pushed through.



Gloves are attached to the glove port via the interchangeable glove rings using a neoprene 7" port diameter 710mm long gauntlet- style glove.



A bag system can also be connected to a glove ring and used to supply food, liquids or medicine to the patient or can be used to remove items from the device (items should not be removed from the AeroPod during transfer of contaminated patients).

Instructions on Use

Changing Gloves

If gloves become compromised they can be replaced as follows:

1. Secure inner port cap.
2. Move the heavy duty band to the inner groove, with the old glove secured over the outer groove without the band.
3. Place the new glove inside the old one and stretch over the old and onto the inner groove.
4. Secure the new glove with the securing band.
5. Once the new glove is firmly secure, while holding the new glove and band in place, gently pull the old glove into the Pod and place in a waste bag (waste should be removed when the Pod is opened at the receiving unit).

Using the Supply Bag

1. Secure inner port cap.
2. Place the item(s) to go into the Pod in the supply bag (if no supply bag, secure that first).
3. Place another supply bag into the first supply bag.
4. Move the securing band to the inner groove, with the first bag remaining secured over the outer groove without the band.
5. Stretch the second bag over the old and onto the inner groove.
6. Secure the second bag with the securing band.
7. Once the second bag is firmly secure, while holding the second bag and securing band in place, gently pull the first supply bag and its contents into the Pod.
8. Place the used supply bag into a waste bag (waste should be removed when the Pod is opened at the receiving unit).

Using Connector Ports

The Connector Ports are made up of five components:

- Main body
- Cap
- Silicone bung
- Oxygen in-line connector
- Cone

The connector ports allow for the connection of IV lines and electrical monitoring equipment. The silicon bungs are designed to create a seal when not in use if the cones are not in place. The pass-through opening is 46.4mm in diameter.

The cones are cut at the end to allow the tubes and wires to pass through and are then sealed through taping round the end.

The in-line oxygen connector can be put in the port and is sealed by screwing on the connector cap. The oxygen connector is filtered preventing back flow of contamination.



Ventilation System

AeroPod offers both positive and negative pressure options. The air ventilation system consists of:

- 2 HEPA filters
- 2 Hoses
- CleanAir2F Ex APR (Air Purifying Respirator) system

The APR unit can provide an adjustable airflow of 120 to 235 litres per minute. Tested according to EN12941 TH2/TH3, EN 12942 TM2/TM3/ IP65.

The APR unit has a full colour LCD that displays key information including:

- Flow rate resistance
- Battery capacity (% remaining)

The unit has an alarm that sounds at critical points:

- High resistance (blocked filters)
- Low output/ flow
- Low battery

The APR unit is powered by a li-ion 14.4v 2.6AH rechargeable battery. Time to fully charge the battery is 3 hours



CleanAir 2F Unit

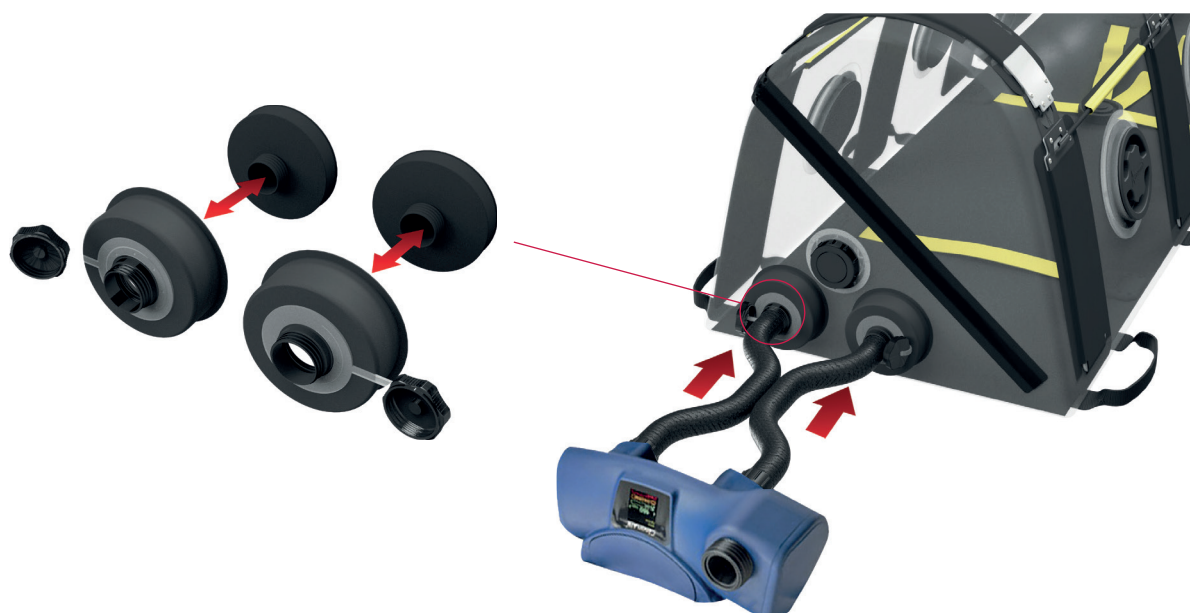
CleanAir P3 Particle Filter

The ventilation works on pressure +/- 21 Pa and provides more than 15 full air exchanges per hour.

Negative Pressure

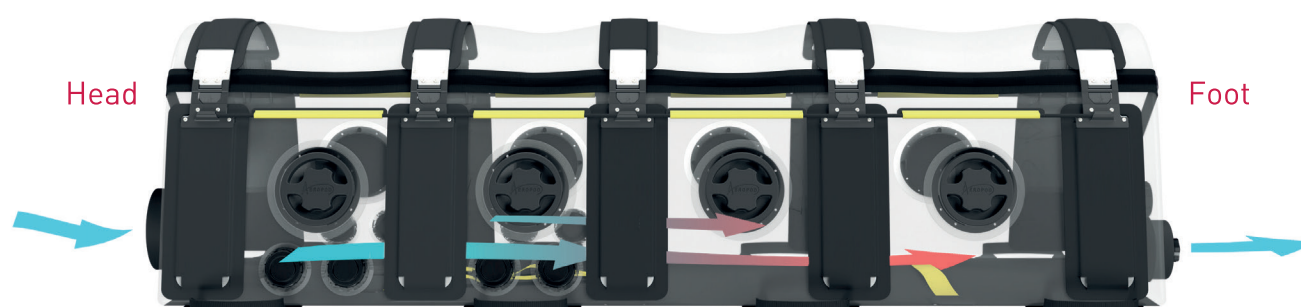
HEPA filters are installed on the inside of the AeroPod and are connected directly to the hoses. The hoses are connected to the APR system. HEPA filters are connected to the other end of the AeroPod allowing inflow of air into the Pod. The CleanAir device should then be switched to negative pressure.

The APR unit is connected to the stretcher or worn around the waist of attending staff.



Negative pressure draws air in through an inlet filter at the head end of the device expelling it through filters at the foot end of the device and out through the APR unit.

This generates negative pressure inside the device, relative to the external environment.

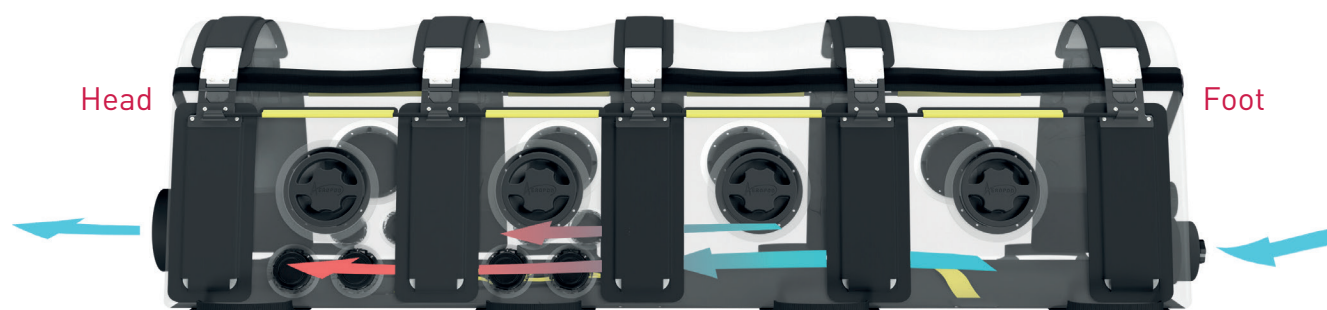


Positive Pressure

Positive pressure supplies air into the device through the APR unit via HEPA filters externally attached to the APR, increasing the pressure inside the device, relative to the external environment. Air is pushed in from the APR system at the foot end of the device and creates an airflow up and over the body and is expelled through the HEPA filter at the head end. The APR unit is then attached to the stretcher or round the waist of a member of staff.



Note: Use positive pressure yellow adaptor.



Assembly Instructions

1. Remove from bag and unroll onto stretcher. Unfold ribs and lay flat. Adjust ribs to ensure material is not trapped in hinges.
2. Attach 6 anchor straps to stretcher using D-Ring fasteners.
3. Insert lateral rod through the yellow sheaths on side of unit.
4. Flop this side of unit over the side of the stretcher.
5. Insert second rod into opposing side yellow sheaths.
6. Insert passive air filter and HEPA filters and attach air purifying respirator (negative pressure for contaminated patients / positive pressure for immunocompromised patients).
7. Transfer patient into AeroPod as per local protocol.
8. Prepare AeroPod by feeding patient monitoring wires through connector port cones, using tape to seal and inserting glove ports ready for use. The netting inside the unit is suitable to house small items that may be needed during transport.
9. Bring side of Pod up over patient while supporting it to stop the sides falling on the patient, and connect all struts starting with 1,3 & 5, ensuring all hinges are locked correctly.
10. Prepare glove and supply ports and apply internal and external caps to seal unit.
11. Fully fasten gas-tight zip and ensure zip is fully closed and buttoned.
12. Ensure airflow is functioning correctly (negative pressure -21pa positive pressure +21pa).
13. Clip lateral rods into lateral rod clips on ribs and feed ends into stoppers within rod clips.

Patient Loading

This process should be undertaken by staff who are familiar with the device and have completed training in its use.

1. Assemble the AeroPod as per the Assembly Instructions up to the point of loading the patient, prior to entering the patient area/ red zone.
2. Assess what equipment is required in the Pod for transfer and take into the patient area/ red zone with Pod.
3. Don (put on) the required level of PPE prior to entry (as per local/ national policy).
4. Take the Pod into the patient area/red zone.
5. Position the Pod up to the patient bed/trolley for transfer into the Pod. Apply brakes.
6. Transfer the patient onto the Pod, as per local protocol. Ensure patient has a small pillow under their head and wedge under their knees for comfort (as appropriate), and the harness is fastened and tightened.
7. Place required equipment into the Pod for transfer, feeding wires/tubes through connector ports, as necessary. **IMPORTANT:** Remember to seal cones with tape after threading wires/tubes through aperture. The netting inside the unit is suitable to house small items that may be needed during transport.
8. Ensure airflow is on.
9. Assemble the Pod around the patient, as per Assembly Instructions, ensuring the patient's face is protected during the process. Insert rods into rod clips on ribs and feed ends into stoppers.
10. Prepare glove and supply ports, and attach internal and external caps. Ensure spare gloves and bags are available.
11. Ensure the airtight zip is fully pulled and buttoned.
12. Ensure airflow is functioning correctly (the envelope should concave slightly when negative pressure is functioning correctly).
13. Complete Assembly Checklist (see page 29) to ensure correct assembly.
14. Once the patient is sealed in the Pod, if moving a biologically contaminated patient, it should be moved to an amber area and the outside of the Pod, airflow system and trolley, cleaned with an appropriate disinfectant tested to EN14885 standard or equivalent, to render it safe for transfer e.g. Peracetic acid, Chlorine dioxide, 1,000ppm hypochlorite solution (10,000ppm hypochlorite solution for body fluid spillages), used as per manufacturer's instructions. Seek further advice for decontamination for a chemically contaminated patient. No film warranty is provided if AeroPod is decontaminated with aggressive solvents or incorrectly diluted cleaning agents
15. Once disinfected the Pod and trolley are safe for movement to a clean area/ green zone and transfer.
16. Staff must then doff (take off) their contaminated PPE before entering a clean area/green zone.
17. Staff can use the glove ports for accessing the inside of the Pod during transfer.

Items should only be put into the Pod via the bag port during transfer. When moving contaminated patients, no waste should be taken out of the Pod until delivery to the receiving unit and the Pod is opened in the patient area/red zone.

NOTE:

- Remove patient shoes where possible.
- Use inflatable under-knee cushion for taller patients.

Patient Unloading

It is essential that the Pod is opened in the order described below to prevent damage.

This process should be undertaken by staff who are familiar with the device and have completed training in its use.

1. The receiving unit staff must don appropriate PPE (as per local / national policy) prior to opening the Pod in the patient area / red zone.
2. Once the Pod / patient is in the patient area/red zone and staff are in PPE the Pod zip can be opened.
3. Once the zip is **FULLY** open, fold side down the side of the trolley and position trolley next to patient bed. Apply brakes.
4. Detach any monitoring equipment that has been fed through the connector ports before patient movement.
5. Unfasten patient harness and transfer patient to patient bed, as per local protocol.
6. Move Pod to decontamination area/amber zone for decontamination.
7. If the contaminated Pod needs to be moved through a clean area / green zone it can either be zipped up and the outside cleaned, as per patient loading section, prior to movement or rolled up and placed in a plastic bag for movement to a decontamination area, remembering to disinfect the outside of the bag prior to transfer out of the patient area/red zone or amber zone.

Decontamination Protocol

The following gives guidance for the decontamination of the AeroPod following the transfer of a patient with biological contamination. Further advice will need to be sought for decontamination following chemical contamination.

Equipment for Decontamination

This is the minimum equipment required to complete the decontamination of the AeroPod:

- Appropriate disinfectant as determined by local / national policy (e.g. Peracetic acid, Chlorine dioxide, 1,000ppm hypochlorite solution (10,000ppm hypochlorite solution for body fluid spillages), tested to EN14885 standard or equivalent, used as per manufacturer's instructions.
- Dry cloths, as necessary.
- Disposable bowl if using a solution.
- Appropriate waste equipment (e.g. category B, category A waste equipment).
- Fumigation device according to local / national policy.
- Screwdriver for unfastening patient harness and anchoring straps. (NOTE The patient harness manufacturer only recommends soap and water for harness cleaning, therefore when disinfection is required the harness will need to be replaced.)

Decontamination Procedure

- Don appropriate level of PPE for equipment decontamination in a contained area.
- Remove any waste, the gloves and waste bag from ports, and all filters from the Pod and dispose of the **patient harness** as per national protocol.
- Disinfect all surfaces of the Pod internally and externally with the appropriate disinfectant, as per national protocol.
- Disinfection should include the following:
 - Patient head and knee pillows (discard if grossly contaminated with body fluids).
 - Internal and external porthole caps once removed.
 - Lateral Rods.
 - APR, including disposal of filters.
 - Pod filters housing.
 - Internal and external envelope.
 - Top Ribs and Side Struts.
 - Underside of Pod.
- Once surface disinfection of the Pod is complete disinfect the environment of the decontamination area / amber zone.
- Staff should then doff PPE, as per national policy.
- For final enhanced decontamination place all components of the Pod in a manner where fumigation / enhanced decontamination treatment can access all surfaces.

- Undertake additional fumigation/enhanced decontamination treatment of the Pod with product as per local / national policy, following manufacturer's instructions. Products that can be used on AeroPod include; Hydrogen peroxide vapour and ozone. UV treatment can also be used as an alternative to fumigation if local / national policy permits. Please contact the company for any further advice on compatibility of other products.

NOTE:

- Once final disinfection is complete the product is ready for reassembly. See reassembly instructions.

Transportation of Contaminated / Used Pod

If unable to completely decontaminate the Pod after patient unloading the Pod should be safely packaged prior to transport for decontamination in another location. Undertake this in a contained zone away from the patient area (this is essential if dealing with an airborne disease). A suggested procedure for this is as follows:

Amber Zone

1. Dispose of all waste and disposable elements and clean visible body fluid spillages. DO NOT remove the harness at this stage.
2. Reseal the unit and decontaminate the outside with an appropriate disinfectant (as discussed earlier in document), keeping the Pod on negative pressure.
3. Remove the lateral rods.
4. Remove the air purifying unit keeping the filters insitu.
5. Unzip and collapse the Pod, with minimal contact with the inside of the Pod as possible.
6. Roll Pod up disinfecting the base as it is rolled.
7. Place in large waste bag (e.g. Category A waste bag or a contaminated mattress bag if yellow waste bag is too small) swan neck and cable tie the bag.
8. Disinfect outside of the bag.
9. Pass to a clean zone/area and staff in clean gloves and an apron should seal the Pod in another bag.

Green / Clean Zone

The Pod should now be ready for transport. Ensure no sharp objects have contact with the bagged Pod to prevent internal exposure.

Attention

AeroPod 15 should be thoroughly cleaned and decontaminated after each use.

All consumable items - gloves, filters, bags must be disposed of as per local / national policy.

Ensure harnesses are properly disinfected.



Emergency Procedures

Blower Failure

If this happens an alternative blower, on-board vehicle system or manual system can be connected in order to maintain the ventilation.

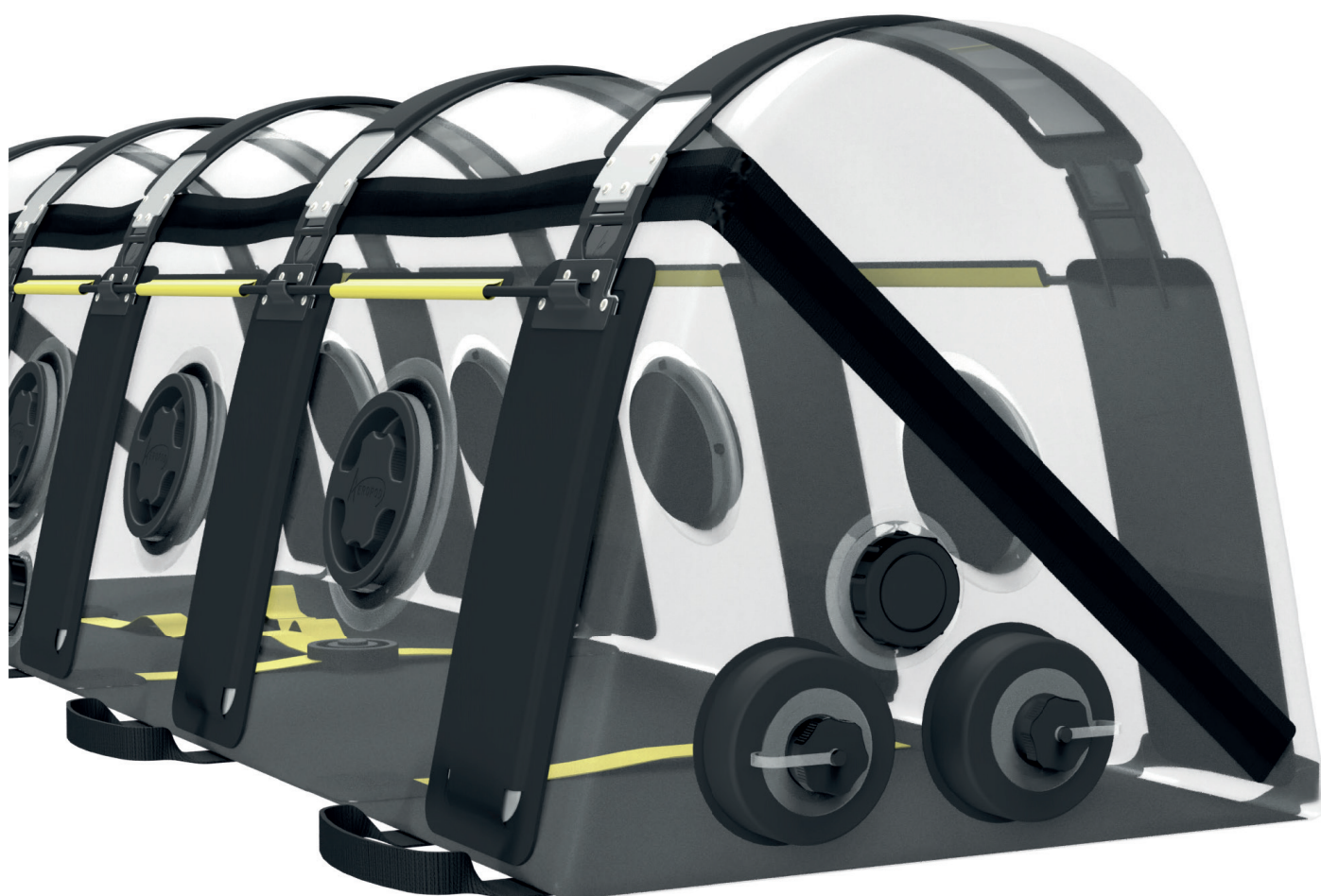
Glove Damage

If gloves are compromised an alternative glove port should be used or the gloves can be replaced using the procedure described on page 13.

Evacuation

The AeroPod zip can be opened at either the head or the foot end and the structure is easily unclipped to allow a quick full exit if necessary.

In an extreme emergency the device can be cut open with scissors.



User Requirements

AeroPod is intended for use by healthcare professionals or first responders only after an appropriate level of training has been completed.

Organisational Requirements

Organisations must ensure they have the appropriate policies and procedures in place prior to using the AeroPod.

This includes:

- Risk assessment processes.
- Safe clean area for storing of equipment.
- Appropriate area for decontamination.
- Regular training and CPD opportunities for users.

BBL Protect are able to provide appropriate training and can assist with the development of in-house policies and procedures.

Attention



AeroPod 15 is designed to reduce and minimise risk, when used correctly.

It does not totally eliminate risk of infection / contamination.

All operators of AeroPod must use appropriate PPE, follow organisational policies and procedures, ensure AeroPod is appropriately decontaminated after use and stored as per manufacturers instructions.

Product Specifications - AeroPod 10

Technical data:

Outer dimensions: L=1860mm W=490mm H=480mm.

Weight:

Patient weight: max 150 kg.

Patient Length: max 198 cm.

Operating temperature between: 0° and +40° C.

Patient retention / safety belts:

Aviation standard quick release 4-point torso / hip restraint, shoulder straps and lower leg strap.

Access ports:

Access ports : 8 ports for gloves,

Waste bag or sluice bag configurations.

Connector ports: 9 ports with cut and seal cones for inlet of IV lines, monitoring cables and similar equipment. Max 46.4 mm opening.

Air system:

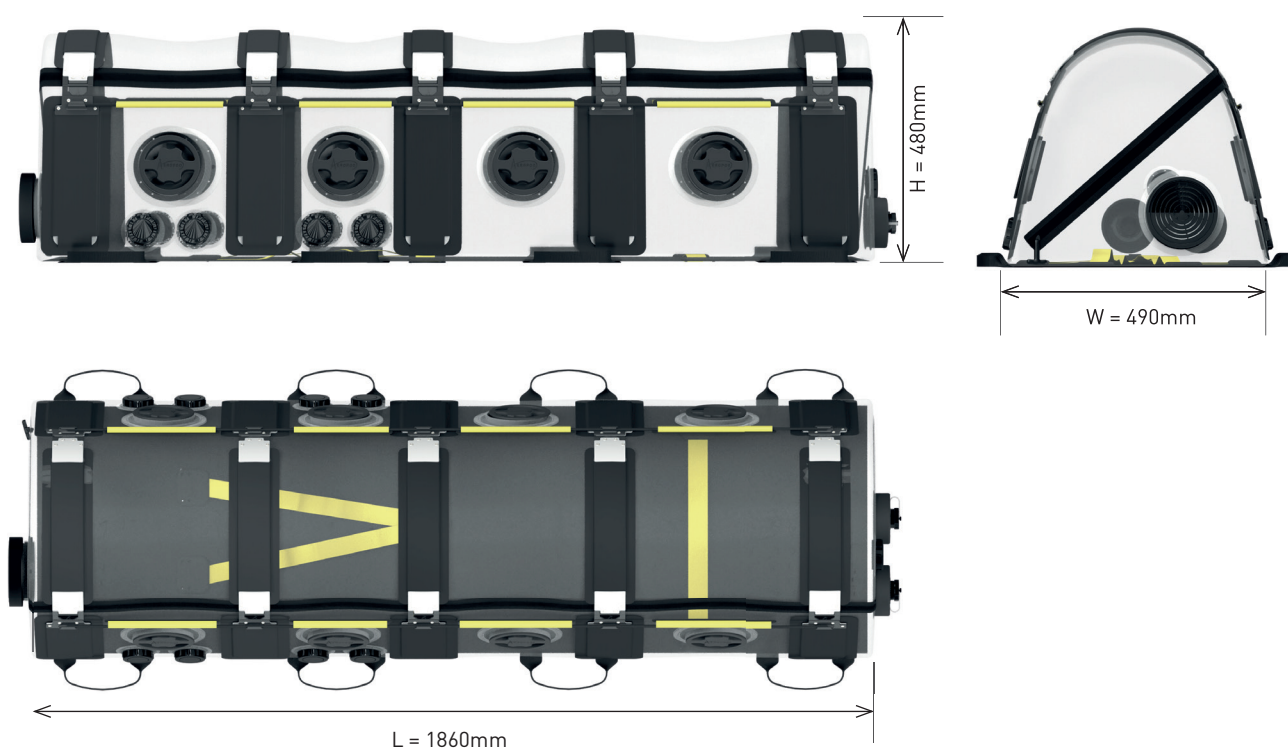
Blower unit: CleanAir Chemical 2F.

Power source : Rechargeable Li-ion battery 14,4V 2,6 Ah, CE1024.

Air exchange : More than 15 air exchanges per hour.

Operating Negative pressure: min 21 Pa relative to the environment.

Operating Positive pressure: min 21 Pa relative to the environment.



Product Specifications - Aeropod 15

Technical data:

Outer dimensions: L=1902mm W=550mm H=523mm.

Weight:

Patient weight: max 150 kg.

Patient Length: max 198 cm.

Operating temperature between: 0° and +40° C.

Patient retention / safety belts:

Aviation standard quick release 4-point torso / hip restraint, shoulder straps and lower leg strap.

Access ports:

Access ports : 8 ports for gloves,

Waste bag or sluice bag configurations.

Connector ports: 9 ports with cut and seal cones for inlet of IV lines, monitoring cables and similar equipment. Max 46.4 mm opening.

Air system:

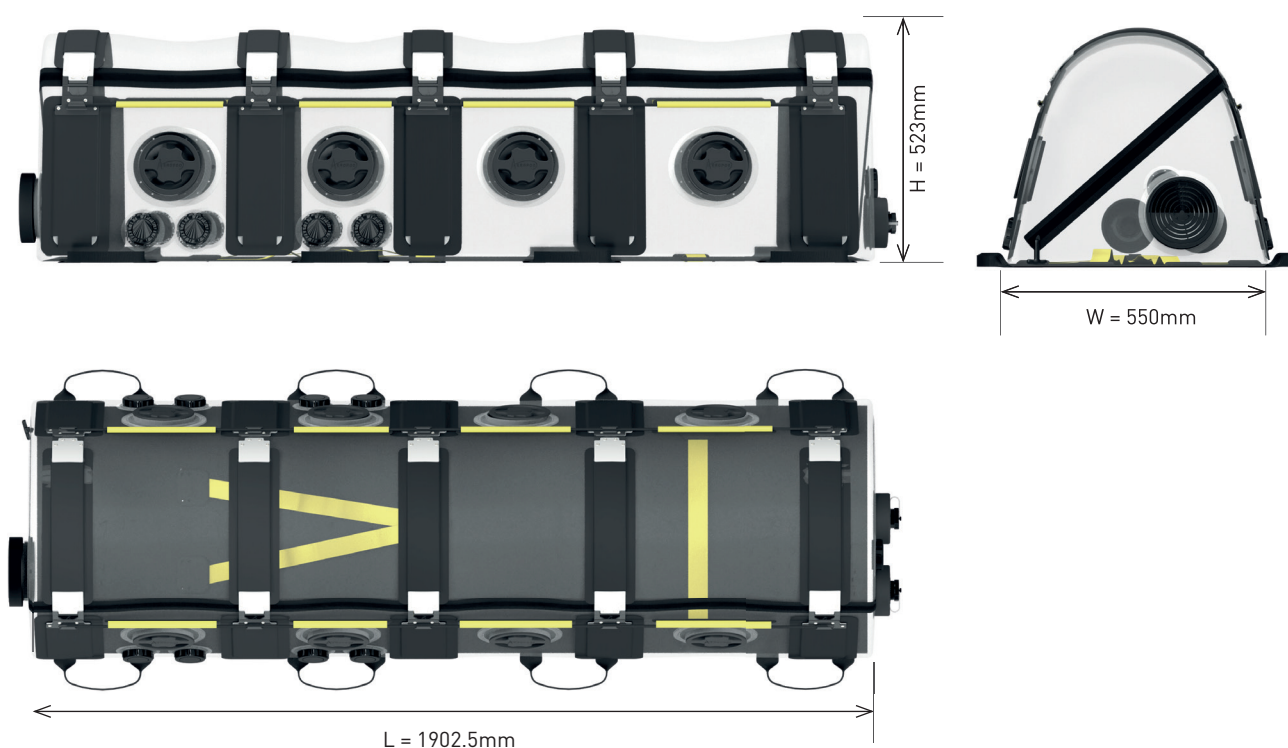
Blower unit: CleanAir Chemical 2F.

Power source : Rechargeable Li-ion battery 14,4V 2,6 Ah, CE1024.

Air exchange : More than 15 air exchanges per hour.

Operating Negative pressure: min 21 Pa relative to the environment.

Operating Positive pressure: min 21 Pa relative to the environment.



Storage of AeroPod

After each use, following decontamination it is important to check AeroPod for any wear or damage that may have occurred during use, including missing or broken parts.

Ensure the Pod is fully dry before packing away to avoid any risk of mould or mildew.

AeroPod should be stored in its own bag in a cool dry environment:

Storage temperature **10 - 20 degrees Celsius.**
 50 - 68 degrees Fahrenheit.

When storing AeroPod ensure the Pod is clear of any sharp objects or debris which may damage the Pod. AeroPod should be checked on a monthly basis using the checklist below.

NB: Please note it is recommended that a requirement to take the AeroPod out of the bag for airing and checking once a month be included into organisational policies and procedures to ensure the Pod has received no damage whilst being stored and is ready for use.

Storage and Checking of AeroPod

AeroPod Integrity Checklist

AeroPod should be checked for the following before storing and on a monthly basis:

- Any cracks or tears in the materials.
- All internal harnesses are working and have no damage.
- There are no biological substances trapped in e.g. harness buckles.
- Anchor straps are complete.
- All double D rings are present.
- All glove port caps and glove rings are present.
- Zip is working with no damaged or missing teeth.
- Top ribs and side struts have no wear and tear.
- Hinges have no damage or cracks.
- Locking pins are not bent.

AEROPOD ASSEMBLY AND PATIENT LOADING CHECKLIST

Element	Completed
Unit checked for trapped hinges on unrolling and prior to assembly.	
6 safety straps attached to stretcher using D-Ring fasteners.	
First structural rod inserted through yellow sheaths on side of unit.	
Side of unit placed over the side of the stretcher.	
Second structural rod inserted into opposing side yellow sheath.	
Passive HEPA air filter and x2 blower HEPA filters inserted and air purifying respirator attached (negative pressure for contaminated patients / positive pressure for immuno-compromised patients).	
Patient's shoes removed.	
Patient harness fastened and tightened.	
Patient pillow and inflatable under-knee cushion insitu – as necessary	
Monitoring wires fed through connector port cones and sealed with duct tape.	
Other additional equipment placed in netting pockets.	
All necessary glove ports and supply ports assembled, with spare gloves and bags available.	
All ports sealed.	
AeroPod assembled locking hinges 1, 3 and 5 first.	
All other hinges locked.	
Gas-tight zip fully fastened and buttoned.	
Airflow functioning correctly (negative pressure -21pa positive pressure +21pa).	
Rods clipped into structural connectors on ribs and ends fed into stoppers.	

Details of person completing checklist

Name (print): _____

Designation: _____

Signature: _____

DECONTAMINATION RECORD

Use no.	Purpose for use	Type of Decontamination completed	Date	Signature	Print Name
1					
2					
3					
4					
5					
6					
7					
8					
9					
10					
11					
12					
13					

DECONTAMINATION RECORD

Use no.	Purpose for use	Type of Decontamination completed	Date	Signature	Print Name
14					
15					
16					
17					
18					
19					
20					
21					
22					
23					
24					
25					



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A detailed 3D rendering of the Aeropod device, a medical transfer aid. It features a grey frame with multiple straps and buckles, and several circular components with the 'AEROPOD' logo. The device is shown from a side-on perspective, highlighting its complex structure and adjustable parts.

AEROPOD®

**The Next Step in
Patient Transfer**

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