

INFANT RESUSCITATION CABINET

OPERATING INSTRUCTIONS



CE0086

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INTRODUCTION

This document contains the instructions for the operation, cleaning and basic maintenance of the wall mounted Viamed Infant Resuscitation Cabinet.

Viamed is not liable for the proper functioning of any part of this product if it is not operated according to the instructions, if the maintenance recommendations in this manual are not followed, or if repairs are carried out using non-approved components.

Only qualified personnel should carry out servicing of this equipment. Maintenance documents can be obtained through your local dealer or direct from Viamed.

Servicing personnel must be aware of the potential clinical implications of incorrectly serviced equipment.

The personnel who work with this equipment should read this manual carefully and should fully understand all instructions contained therein. The manual should be kept so that it can be easily inspected; it is advisable to store it in an easily accessible place.

If any function or part of this manual is not clear, please contact Viamed or your distributor in order to obtain further information or clarification.

Overview

The Viamed Infant Resuscitation Cabinet contains a carefully selected group of products; each main item can be purchased as a single item for use individually.

Viamed have combined these devices into one product in an arrangement ideally suited to Maternity and Neonatal Intensive Care environments.

Devices forming component parts of the Resuscitation Cabinet:

- Cabinet with drop down door, integral bed and mattress
- Wall-mounted Ceratherm 600-2 radiant warmer
- Tom Thumb infant resuscitation unit
- Low suction controller
- Suction receiving jar
- Digital Apgar timer
- Storage bins

All the components carry the correct and relevant CE marking.

WALL MOUNTED RADIANT WARMER



The radiant warmer is mounted on a double-hinged arm, allowing the unit to be folded flat to the wall when not in use.

Precautions

The radiant warmer must not be used in the immediate vicinity of explosive anaesthetic gases or mixtures.

The distance between the surface on which the patient lies, and the lower edge of the radiant warmer, must not be less than 80cm. If this instruction is not followed, prolonged exposure to the heat radiation may cause burns.

“Danger of burns”: Contact with the protective guards, element and reflector should be avoided.

“Danger of fire”: The grille on the upper side of the radiant warmer must remain unobstructed at all times.

Do not place any items on the grille.

Patients must not be left unattended whilst the radiant warmer is in use.

Do not operate the device if the casing has been removed or is damaged.

Servicing and maintenance should be restricted to qualified technical personnel.

Use of the radiant warmer should be restricted to trained medical personnel.

Use of a radiant warmer can increase fluid loss from a patient.

Handle the warmer with care; the ceramic heating element is fragile.

Do not touch the heating element at any time as this can adversely affect the performance of the element.

The heating element gets very hot in use; take care if the warmer has recently been used or tested, and allow it to cool before storage. The side handles remain cool to touch.

Each Ceratherm 600-2 Infant Radiant Warmer is safety tested to Class 1, Type B and a test report is supplied. It is recommended that the equipment be retested for safety and function upon receipt, and at least annually thereafter.



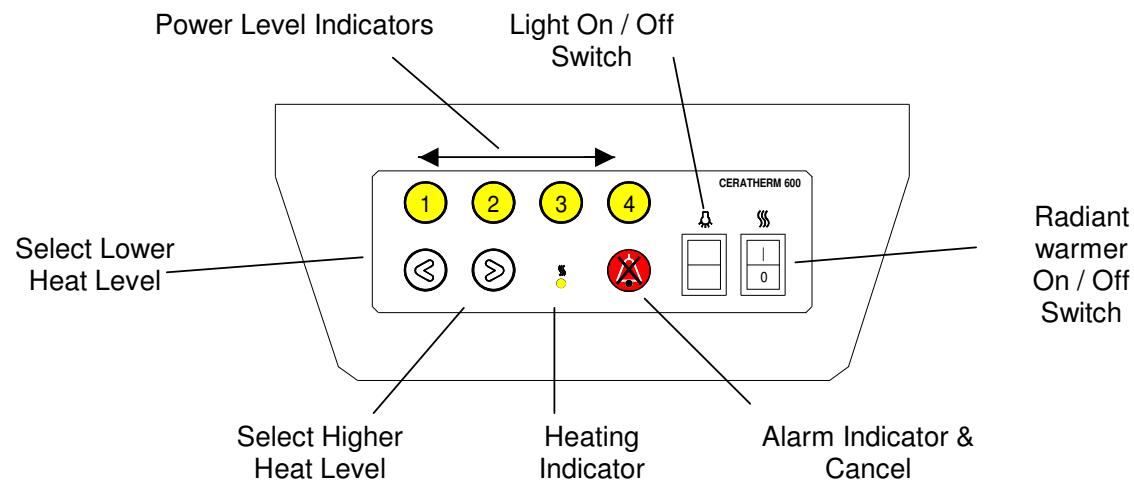
WARNING!

The radiant warmer gets hot whilst in use.

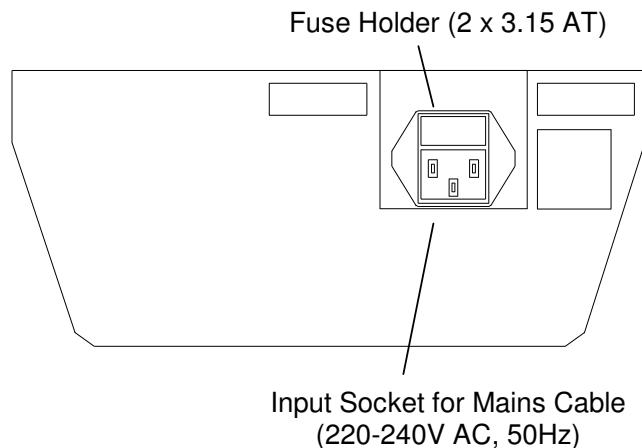
Do not obstruct the upper ventilation grille.

Use only the side handles, which remain cool, to reposition the unit.

Front



Rear



Adjusting the Heat Output

Switch radiant warmer On/Off switch to ON. This switch is the master power switch for the device; no electrical functions are active with this switch in the OFF position.

Using the forward and reverse buttons, < and >, set the desired heat output level. The current heat output setting is shown by an illuminated yellow indicator ①, ②, ③ or ④.

The heat output can be adjusted using the four heat output settings (① - ④). The factory-set levels corresponds to the following values, the values indicated correspond to a percentage of the maximum power output of the heating element:

Level 1	= 25%	①	②	③	④
Level 2	= 50%	①	②	③	④
Level 3	= 75%	①	②	③	④
Level 4	= 99%	①	②	③	④

The various power levels can be used as follows:

Level ①: To keep the support surface warm for continuous operation.

Level ②: For normal operation, on changing and examination areas.

Level ③: Additional warmth for resuscitation, the labour room or for the operating theatre.

Level ④: For increased heat, in the operating theatre, during anaesthesia, or for adults.

Using the Halogen Lamp

The non-dazzling halogen lamp is used for illuminating the patient area.

To turn on the lamp, set the light On/Off switch to ON. Note: the lamp can only be used when the radiant warmer On/Off switch is in the ON position.

If it is required, the halogen lamp can be used without the heating element, to do this adjust the heat output of the radiant warmer to zero. The heat output is set to zero when the reverse button, <, is repeatedly pressed until all yellow indicators (① - ④) are extinguished.

LED Heating Indicator

The Heating Indicator illuminates when the element is being heated, and is off when it is not. This 'duty-cycle' occurs over a period of a second, the percentage of time that the indicator is illuminated during that second represents the power level in terms of a percentage of the maximum output.

Alarm and Safety-Power Function

The radiant warmer incorporates a safety feature, which reduces the output power if the device is left unattended. An audible and visual alarm is triggered a period of 15 minutes after the last key-press; this is to inform the operator that the device is still active.

The safety alarm activates an audible alarm signal, which consists of a series of beeps over a 5 second period, and the red Alarm Indicator on the front panel begins to flash.

The alarm is cancelled by pressing the flashing Alarm Indicator, if this is done during the first 8 seconds after the onset of the alarm condition, the radiant warmer will continue to function at the current power setting, the Alarm Indicator stops flashing, and the 15-minute timer begins again.

If, after 8 seconds from the onset of the alarm condition the alarm has not been cancelled, the heat output of the radiant warmer reduces to a pre-set value, which is 20% of the maximum power output of the device.

Note: when the radiant warmer enters the reduced power setting, the yellow power level indicator remains illuminated at the previous power setting, this is to inform the user which power setting that the device will return to when the alarm is cancelled.

When the alarm is cancelled, the radiant warmer reverts back to the previous power setting, the Alarm Indicator stops flashing, and the 15-minute timer begins again.



WARNING!

The radiant warmer must not be left unattended whilst in use.

RESUSCITATION CABINET



The cabinet has been designed to conceal the internal products when closed, and provide easy access when open.

The drop down platform has an integral bed, on which the mattress is placed. Additional blanket material may be placed on the mattress, but care should be taken to ensure that this does not become trapped in the space between the platform and the cabinet when raising the platform.

Excessive weight should not be applied to the drop down platform. The maximum loading, including the pressure applied by the operator(s), should be no greater than 25 Kg.

Whilst in use, the patient should be under close observation at all times.

When closing the cabinet, ensure that there are no obstructions; the cabinet should close easily and without significant force. Once the door is in the upright position, turn the catch to secure it in the closed position.



CAUTION

The door should not present resistance or need to be held closed in order to secure the catch.

Securing the catch whilst forcing the door closed may cause damage to the catch, damage of this type is not covered by the warranty.

LOW SUCTION CONTROLLER

The Viamed Infant Resuscitation Cabinet is supplied with one of two low suction controllers, designed specifically for medical use: the Oxylitre S714 or the Therapy Equipment 4725-3M.

Vacuum Source Connector

The suction controller is supplied with a 3m vacuum hose assembly, which terminates in a British Standard (BS5682) probe. The device is intended for connection to a wall vacuum outlet.

Patient Inlet Connection

The inlet connection forms an integral part of a detachable filter cartridge or Disposable Hydrophobic Filtration Unit. This filtration unit has been designed to prevent the ingress of fluids into the controller and the pipeline system.

Once the filter has been used for patient therapy, or if wetted for any reason, it must be replaced. If the filter membrane is wet, the chemical coating on it will prevent fluid passing through it and may restrict the flow of suction.

Vacuum Gauge

The controller is fitted with an easy to read, dual scale colour coded gauge.

Model	Scale readings
Oxylitre S714	0 to 200 mmHg (0 to 25kpa)
Therapy Equipment 4725-3M	0 to 300 mmHg (0 to 40kpa)

Safety Valve

The suction controller is fitted with an internal safety valve system. This will protect the suction controller from being damaged if inadvertently connected to a positive pressure outlet.

Setting Suction Levels: Oxylitre

To increase the level of suction, rotate the control knob on the top of the suction controller in an anti-clockwise direction.

To decrease the level of suction, rotate the control knob on the top of the suction controller in a clockwise direction.

The Oxylitre S714 suction controller does not have a separate ON/OFF control, the device is turned off by decreasing the suction level to the minimum setting.



WARNING

The suction level must be set before each use to a level determined by guidelines within the customer's own organisation.

Setting Suction Levels: Therapy Equipment

To increase the level of suction, rotate the control knob on the top of the suction controller in a clockwise direction.

To decrease the level of suction, rotate the control knob on the top of the suction controller in an anti-clockwise direction.

The Therapy Equipment 4725-3M has a separate ON/OFF control in the form of a 'flag' lever on the right hand side of the device.

To turn the suction on move the lever to the vertically downward position, the word 'ON' is displayed on the flag. To turn the suction off move the lever to the vertically upright position, the word 'OFF' is displayed on the flag.



WARNING

The suction level on the Therapy Equipment low suction controller will remain set at the level last used.

The suction level must be set before each use to a level determined by guidelines within the customer's own organisation.

SUCTION RECEIVING EQUIPMENT

The Viamed Infant Resuscitation Cabinet is supplied with the VacSax disposable suction receiving system as standard, or by request, a reusable Therapy Equipment suction jar.

Viamed do not supply suction catheters, however, the system is designed for use with standard 10mm suction tubing compatible with a wide range of commonly used suction catheters.

Whichever receiving system is used, ensure that the suction tubing and catheter is replaced after each use.

Instructions for Use; Vacsax System

Ensure jar is clean, wipe inner rim with damp cloth if necessary. If jar is contaminated it may be autoclaved at 121° C or cleaned with water-based disinfectants; see instructions later in this document.

DO NOT USE CANISTER IF IT IS CRACKED.

Remove liner from packaging. If collapsed pull straight. Place liner in jar and push down firmly. Ensure patient port is facing towards 'VacSax' mark on jar.

Connect suction source to liner using taper connector, push into vacuum port using a twisting motion.

Always use a taper connector (Pt No. 3833-007) on vacuum source tubing when using VacSax liners.

Connect patient tubing to patient port. Push patient tubing connector firmly over patient port. *Always use tubing that is a good fit over the patient port.*

Turn on suction and set desired vacuum level. Confirm suction is present at patient end. *Check operation of the rest of suction system.*

If liner has been accidentally over filled, denoted by vacuum cut off and fluid visible in the patient tubing.

- I. Do not turn off suction.
- II. Remove patient tubing.
- III. Push small stopper firmly over patient port.
- IV. Turn off suction and remove taper connector from vacuum port.
- V. Plug vacuum port using long stopper.

Release yellow stopper from its recess in the cap, twist stopper to break apart.

Push stopper firmly over patient port and into vacuum port. *The larger taper plug pushes into the vacuum port, the smaller plug pushes over the patient port.*

Using handles lift liner from jar and dispose of in accordance with hospital policy.

DO NOT THROW AWAY THE TAPERED VACUUM PORT CONNECTOR. Taper may be cleaned using disinfectants or autoclaved at 121°C.

Instructions for Use; Therapy Equipment System

The Therapy equipment suction jar is a reusable jar manufactured from polysulphone, it may be autoclaved at 121° C or cleaned with water-based disinfectants in accordance with the guidelines in the MAC Manual “Part 2; Cleaning (manual) - non-immersion”; see instructions later in this document. Additionally it may be pre-washed using an automated disinfecter operating at 90° C.

Connect the suction controller to the receiving jar using 10mm suction tubing, ensure the tubing is connected firmly onto the connection marked ‘vacuum’.

Connect the patient tubing to the receiving jar using 10mm suction tubing, ensure the tubing is connected firmly onto the connection marked ‘patient’.

Always use tubing that is a good fit over the connectors of the suction jar.

Always clean the jar thoroughly after use.

The jar cradle requires only external cleaning, which must be done in accordance with the guidelines in the MAC Manual “Part 2; Cleaning (manual) - non-immersion”; see instructions later in this document.

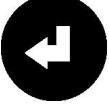
APGAR TIMER

Description

The Viamed Apgar Timer is a battery powered digital timer with preset audible timing indications at 1, 5 and 10-minute intervals.

The timer has a large, clear LCD display and is operated using the keys on the front of the instrument.

The Functions of the Keys

Key	Function
 START	<ul style="list-style-type: none">Starts counting from 00:00.Returns to <i>Counting Mode</i> whilst in <i>Freeze Mode</i>.
 FREEZE/ UNFREEZE	<ul style="list-style-type: none">Enters and exits <i>Freeze Mode</i>. Display 'freezes' whilst the actual elapsed time continues counting, but is not displayed. Exiting <i>Freeze Mode</i> returns to <i>Counting Mode</i>, resuming from the ongoing total elapsed time.
 RESET	<ul style="list-style-type: none">Resets counter, returning timer to <i>Standby Mode</i>.

Standby Mode

The Apgar Timer does not have an ON/OFF switch. When not counting, the Apgar Timer enters *Standby Mode*.

In *Standby Mode* the display is blank with the exception of a flashing colon (:) to indicate the timer is functioning correctly.

If the display is completely blank with no flashing colon, the timer may require battery replacement. Replace batteries on an annual basis.

Counting Mode

To enter *Counting Mode*, press the START button.

When the Apgar Timer begins counting, the display counts upwards from 00:00 in 1-second intervals up to a maximum of 1 hour (59:59 on display), it will then enter *Standby Mode* automatically.

Whilst counting, the timer will indicate the Apgar scoring intervals with audible beeps:

1 minute:	1 beep
5 minutes:	2 beeps
10 minutes:	3 beeps

Freeze Mode

It is possible to freeze the display whilst the actual time continues to count in the background; this is to allow for recording the time from the start of the counting cycle of a particular event.

To enter *Freeze Mode*, press the FREEZE button.

Whilst the display is frozen, the colon (:) continues to flash.

Freeze Mode can be exited and *Counting Mode* resumed by pressing the START or FREEZE buttons.

Upon unfreezing, the timer will continue counting, resuming from the ongoing total elapsed time.

Should an Apgar Scoring interval be reached whilst the display is frozen, the timer will still beep to indicate that interval, but will remain in *Freeze Mode* until exited by the user.

Reset Function

The *Reset Function* allows the timer to be reset and placed back into *Standby Mode*.

Press the RESET button to reset the timer.

Specifications

Dimensions (excluding bracket)	142 mm (W) x 130 mm (H) X 40 mm (D)
Weight	551 g
Display	4 digit LCD. Digit height: 25mm
Power requirements	4 x AA/LR6/MN1500 1.5 V Alkaline Batteries
Controls	Start, Freeze/Unfreeze, Reset
Audio indications	1, 5, 10 minutes

Battery Replacement

For optimal performance, batteries should be replaced on an annual basis.

If the Apgar timer is to be removed from service or stored, the batteries must be removed.

The LCD flashing between the messages “Lo” and “Batt” indicates a low battery condition, when this message appears, the batteries should be replaced at the earliest opportunity.

To replace the batteries, remove the mounting bracket from the rear of the timer using a 3/16th inch Allen Key, then using a narrow blade screwdriver release the battery cover. Remove the existing batteries and insert new ones (4 x AA/LR6/MN1500 1.5V Alkaline), observing the correct polarity.

Depleted batteries should be disposed of as per local regulations.

Routine Maintenance

It is recommended that the Apgar Timer be checked at least annually to ensure correct performance.

Perform service checks as detailed in accompanying Service Sheet. There are no user serviceable parts inside: if unit does not meet the specification please return to Viamed for repair.

Optional Accessories

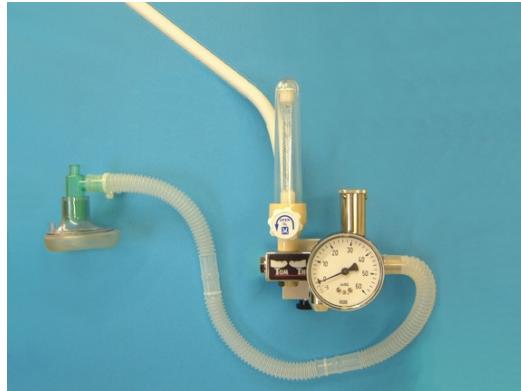
Part Number	Description
0320200	Medirail mounting bracket
0121199	Swivel mount
0121200	Universal mounting clamp

Cleaning and Decontamination

For routine cleaning Apgar timer can be cleaned with isopropyl alcohol, or with a mild detergent solution in accordance with the detergent manufacturer's recommendations.

For decontamination, clean in accordance with the guidelines in the MAC Manual “Part 2; Cleaning (manual) - non-immersion”.

TOM THUMB INFANT RESUSCITATOR



Instructions for Use are attached to the Tom Thumb within the resuscitation cabinet. Please refer to these before using the Tom Thumb.

The Tom Thumb Infant Resuscitator is supplied in one of two variants: the TT490-15, which has a 3m oxygen hose terminated with a BS probe, or (less commonly) the TT480, which is designed to be connected to an existing flowmeter via narrow bore tubing.

The Tom Thumb allows the delivery of oxygen at controlled flow and pressure. A flowmeter is used to control the flow rate of oxygen, and the adjustable pressure relief valve is used to set the maximum pressure delivered.

The Tom Thumb incorporates a secondary precision blow-off valve for added safety.



WARNING

Do not operate the Tom Thumb TT490-15 at flows greater than 15 Litres/min.
Do not operate the Tom Thumb TT480 at flows greater than 10 Litres/min.

Pre-use Checks

Uncap the T piece port.

Adjust the flowmeter to minimum (fully clockwise) and the adjustable valve control to minimum (fully counter clockwise).

Connect the TT490-15 oxygen hose to the external wall oxygen supply. If using the TT480 connect the Tom Thumb to an external flowmeter using narrow bore tubing.

Check that the pressure gauge reads zero, if the needle is outside of the black band then the Tom Thumb requires servicing.

Connect the patient tubing to the Tom Thumb outlet but do not apply the mask to the patient.

Set the flowmeter to the required flow rate.

Occlude the mask and the T piece port, ensuring a completely gas-tight seal.

Gradually turn the adjustable valve control clockwise until the required outlet pressure is shown on the pressure gauge.

Note: when setting the maximum pressure, the needle on the gauge of the Tom Thumb should be stable and oxygen can be heard hissing from the adjustable valve. If the needle is not stable and no hissing can be heard, it is likely that the mask is not entirely occluded; check the seal by repositioning the mask until a stable reading is achieved.

The Tom Thumb is now ready for use.



CAUTION

For use by qualified and trained personnel only.

Use flow rates within the range of the flowmeter.

Adjust outlet pressure after altering the flow rate, if the flowrate is subsequently altered, always re-set the pressure.

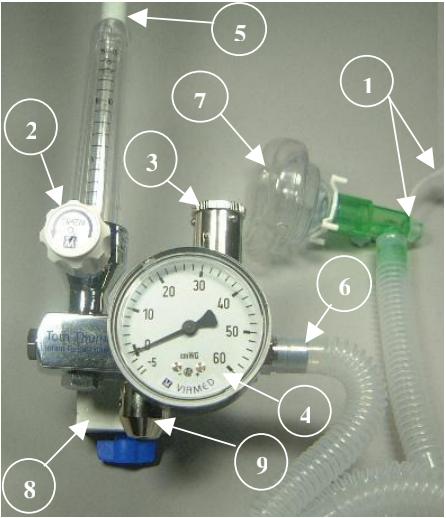
Do not attempt to adjust the safety valve.

Recommended O₂ inlet pressure of 4 bar.

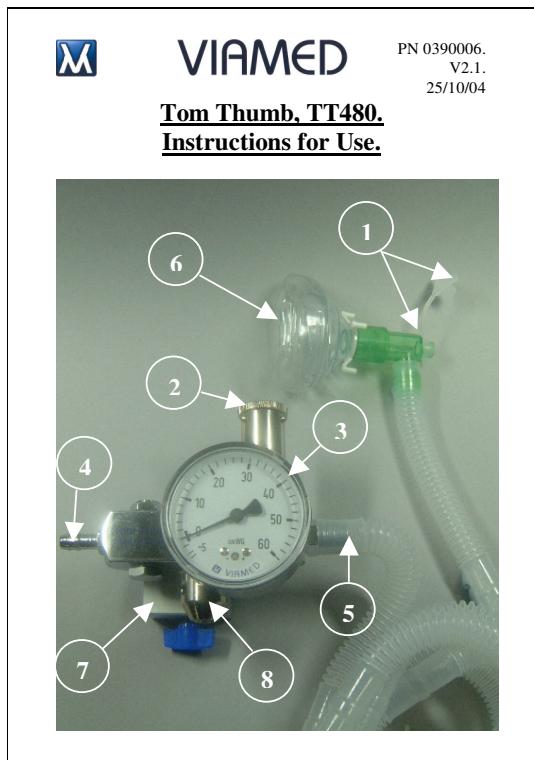
The following pages contain the instruction cards for both variants of the Tom Thumb Infant Resuscitator.

A set of the appropriate instructions is provided with each instrument. Should additional copies be required, they can be created by copying and laminating the relevant instruction page or, alternatively, additional copies can be requested from Viamed.

TT490-15 Instruction Card Template

<p>VIAMED</p> <p>Tom Thumb, TT490-15</p> <p>Instructions for Use</p>  <p>PN 0390007. V2.1. 25/10/04</p>	<p>Pre-use Checks.</p> <ul style="list-style-type: none">Uncap the T piece port ①.Adjust the flowmeter ② to minimum (fully clockwise) and the adjustable valve control ③ to minimum (fully counter clockwise).Connect the flowmeter O₂ hose ⑤ to the external O₂ supply.Check that the pressure gauge ④ reads zero. If not, the Tom Thumb requires servicing.Connect the patient tubing ⑥ to the Tom Thumb outlet but <u>do not</u> apply the mask ⑦ to the patient.Set the flowmeter to the required flow rate.Occlude the mask and the T piece port. Gradually turn the adjustable valve control clockwise until the required outlet pressure is shown on the pressure gauge. <p>The Tom Thumb is now ready for use.</p> <p>Important.</p> <p>For use by qualified and trained personnel only. Use flow rates within the range of the flowmeter. Adjust outlet pressure after altering the flow rate. Do not attempt to adjust the safety valve ⑨. Recommended O₂ inlet pressure of 4 bar.</p>
<p>Guideline for Use during Resuscitation.</p> <ol style="list-style-type: none">Follow the pre-use checks and set the required flow rate and outlet pressure.Apply the mask to the patient and cover the T piece port to inflate the patients' lungs at the set flow rate and pressure.Uncover the T piece port and allow the patients lungs to deflate.Repeat steps 2 & 3 as necessary during the resuscitation of the patient (follow the hospital protocol for resuscitation). <p>Care, Cleaning, Location and Sterilisation.</p> <p>Clean using a damp cloth. The Tom Thumb is not intended to be sterilised. Do not autoclave. Do not allow moisture or foreign matter to enter the safety valve or adjustable valve. Damage will occur if the Tom Thumb is subjected to severe mechanical shock or dropped. The Tom Thumb should be serviced every 12 months, if the pressure gauge does not read zero (outside of the black band) with no flow or if the units' accuracy is doubted. The Oxygen Hoses should be checked every 3 months, and replaced every 4 years as a minimum. The rail bracket ⑧ is designed to fit most medical rails. It is advised that the Tom Thumb is not mounted close to a wall or to the side of an incubator particularly if the gauge is fitted to the end of the body (specials only).</p>	<p>Warranty.</p> <p>Viamed warranty ensures that goods are free from defects of manufacture for a period of one year from the date of shipment from Viamed.</p> <p>Liability shall be limited solely to the replacement and repair of the goods and shall not include shipping costs or other incidental damages.</p> <p>This warranty is null and void if any items are subjected to misuse, negligence, accident, or repairs other than those performed by Viamed or an authorised service centre.</p> <p>Viamed Ltd. 15 Station Road, Cross Hills, Keighley, West Yorkshire, BD20 7DT, United Kingdom. Tel: +44 (0)1535 634542. Fax: +44 (0)1535 635582. Email: info@viamed.co.uk Website: www.viamed.co.uk</p> <p>© Copyright 2003. Viamed Ltd. All Rights Reserved</p>

TT480 Instruction Card Template



Pre-use Checks.

- Uncap the T piece port ①.
- Adjust the external flowmeter to minimum and the adjustable valve control ② to minimum (fully counter clockwise).
- Connect the inlet ④ to the external flowmeter.
- Check that the pressure gauge ③ reads zero. If not, the Tom Thumb requires servicing.
- Connect the patient tubing ⑤ to the Tom Thumb outlet but do not apply the mask ⑥ to the patient.
- **Set the flowmeter to the required flow rate, up to the maximum of 10 Lpm.**
- Occlude the mask and the T piece port. Gradually turn the adjustable valve control clockwise until the required outlet pressure is shown on the pressure gauge.
- The Tom Thumb is now ready for use.



Important.

For use by qualified and trained personnel only.
Use flow rates within the range of the flowmeter.
Adjust outlet pressure after altering the flow rate.
Do not attempt to adjust the safety valve ⑧.
Recommended O₂ inlet pressure of 4 bar.

Guidelines for Use During Resuscitation.

1. Follow the pre-use checks and set the required flow rate and outlet pressure.
2. Apply the mask to the patient and cover the T piece port to inflate the patients' lungs at the set flow rate and pressure.
3. Uncover the T piece port and allow the patients lungs to deflate.
4. Repeat steps 2 & 3 as necessary during the resuscitation of the patient (follow the hospital protocol for resuscitation).

Care, Cleaning, Location and Sterilisation.

Clean using a damp cloth. The Tom Thumb is not intended to be sterilised. Do not autoclave. Do not allow moisture or foreign matter to enter the safety valve or adjustable valve. Damage will occur if the Tom Thumb is subjected to severe mechanical shock or dropped. The Tom Thumb should be serviced every 12 months, if the pressure gauge does not read zero (outside of the black band) with no flow or if the units' accuracy is doubted.

The Oxygen Hoses should be checked every 3 months, and replaced every 4 years as a minimum. The rail bracket ⑦ is designed to fit most medical rails. It is advised that the Tom Thumb is not mounted close to a wall or to the side of an incubator particularly if the gauge is fitted to the end of the body (specials only).

Warranty.

Viamed warranty ensures that goods are free from defects of manufacture for a period of one year from the date of shipment from Viamed.

Liability shall be limited solely to the replacement and repair of the goods and shall not include shipping costs or other incidental damages.

This warranty is null and void if any items are subjected to misuse, negligence, accident, or repairs other than those performed by Viamed or an authorised service centre.

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SERVICING

Service contracts are available; please contact Viamed for further information.

Service the individual items of equipment using their respective Service or Operator's manuals.

For versions of the Tom Thumb that have an integral flowmeter and hose, the manufacturer recommends that the hose is inspected for wear or damage every 3 months, and replaced every 4 years regardless of condition of the hose.

If the Viamed Infant Resuscitation Cabinet has been supplied with a Therapy Equipment Low Suction Controller, the manufacturer recommends that the vacuum hose be replaced at least every 4 years, regardless of the condition of the hose.

If the Viamed Infant Resuscitation Cabinet has been supplied with an Oxylitre Low Suction Controller, the manufacturer recommends that the vacuum hose be replaced at least every 5 years, regardless of the condition of the hose.

CONSUMABLES AND PARTS LIST

Note: the Viamed resuscitation cabinet is available in different configurations, the parts list contains equipment from all variants; care must be taken when re-ordering replacement parts. If in doubt, please contact Viamed to determine the correct item.

Parts	
Part no.	Description
0310002	Complete resuscitation cabinet including all parts
0310025	Wall mounted radiant warmer
0330120	Element for radiant warmer – 600W ceramic
0330125	Halogen lamp for radiant warmer, 20W
0330105	Cabinet body with shelf and medirail
0330009	Resuscitation cabinet door catch
0320020	Resuscitation bed, 35cm (14") wide
0320043	Mattress, 35cm (14") wide
0310035	Low suction controller (Oxylitre S714)
0330040	Suction hose assembly, 3m (Oxylitre)
0310037	Low suction controller (Therapy Equipment 4725-3M)
0330033	Suction hose assembly, 3m (Therapy Equipment)
3833-132	VacSax reusable suction canister
3833-007	VacSax tapered suction connectors (pack/10)
0320010	Universal rail clamp with 'V' for mounting Vacsax suction jar
0310100	Apgar Timer (note: mounting bracket sold separately)
0320200	Mounting bracket with rail clamp for mounting Apgar timer
0310020	Size 3 storage box
0310030	Tom Thumb resuscitation unit, type TT480 without flowmeter
0310034	Tom Thumb resuscitation unit, type TT490-15-3M with flowmeter
0330218	Oxygen hose assembly for Tom Thumb, 3m

Consumables	
Part no.	Description
0310104	Filter for Oxylitre low suction controller, single unit
0310105	Filters for Oxylitre low suction controller, box of 30
0310109	Filter for Therapy Equipment low suction controller, single unit
0310110	Filters for Therapy Equipment low suction controller, box of 10
9910-340	VacSax disposable liners (box/25)
0330035	Suction clear tubing, 1m
0330040	Standard patient circuit, single patient use (includes cushion mask)
3210010	NeoPEEP circuit with variable PEEP, single patient use (without mask) – Pink
3210011	NeoPEEP circuit with variable PEEP, single patient use (without mask) – Blue
3210070	Single use silicone face mask, size 00
3210071	Single use silicone face mask, size 01
3210072	Single use silicone face mask, size 02
0120140	Silicone adapter for connecting NeoPEEP circuit to Tom Thumb (note: required for NeoPEEP circuit, but not for the standard patient circuit)
Operator and Service Manuals	
Part no.	Description
0391045	Resuscitation cabinet operating instructions
0391050	Resuscitation cabinet installation instructions
0390007	Tom Thumb TT490-15 operating instruction card
0390006	Tom Thumb TT480 operating instruction card
0390022	TT480 service manual
0390023	TT490-15 service manual
0390021	Tom Thumb valves service manual (common to all Tom Thumb variants)

		Document ref: 0391055
		Version: 1.1
Viamed Infant Resuscitation Cabinet Service Sheet		
Hospital / Organisation:		
Location (department, room number):		
Engineer (print):		
Service Date:		
Action	Further action required	OK
Cabinet Body – Serial number:		
Check for overall signs of damage	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check door catch – swivel catch on door	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check door catch – retaining clasp on cabinet	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check integrity of welds on the door hinges	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check hinges for damage or deformation	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check the restraining bars for damage or deformation	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check the cabinet is securely fixed to wall	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check medirails and tighten any loose rails	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check grommets on the hose inlets are in place and intact	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check the bumper stops that prevent door banging during closing	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check 'caution' sticker is in place on the platform	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check bed for damage i.e. cracks, chips	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check all the bed mounting screws are in place	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check mattress for damage i.e. tears, split seams	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Tom Thumb – Serial Number:		
Follow service procedure as detailed in the Tom Thumb service manual	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check oxygen hose, if present, for wear, and replace if necessary (at least every 4 years)	<input type="checkbox"/>	<input type="checkbox"/>
Hose replaced? Yes <input type="checkbox"/> No <input type="checkbox"/>		
Date of last hose replacement:		
Comments:		

Action	Further action required	OK
Low Suction Controller – Serial number:		
Follow service procedure as detailed in suction controller operator's or service manual	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Change filter assembly	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check vacuum hose for wear, and replace if necessary (at least every 4 years)	<input type="checkbox"/>	<input type="checkbox"/>
Hose replaced? Yes <input type="checkbox"/> No <input type="checkbox"/>		
Date of last hose replacement:		
Comments:		
Check rail clamp is securely attached to suction controller	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Replace suction tubing	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Replace tapered connector	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check receiving canister for damage, replace if necessary	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Radiant Warmer – Serial number:		
Note: if the unit is hard wired, a PAT test may not be applicable: consult your organisation's regulations.		
Is unit hard-wired? Yes <input type="checkbox"/> No <input type="checkbox"/>		
PAT test performed? Yes <input type="checkbox"/> No <input type="checkbox"/> Pass <input type="checkbox"/> Fail <input type="checkbox"/>		
Comments:		
Follow service procedure as detailed in radiant warmer service manual	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
APGAR Timer – Serial number:		
Change batteries	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Follow service procedure as detailed in Apgar timer operator's manual	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
General		
Check storage bins present	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Tidy any trailing hoses, using tie-wraps if necessary	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Check Operator's Manuals are in place	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Clean cabinet with isopropyl alcohol	<input type="checkbox"/>	<input type="checkbox"/>
Comments:		
Additional notes or comments:		

CLEANING

Cleaning guidelines in this manual complement those given in the Medical Devices Agency document "Sterilization, disinfection and cleaning of medical equipment: guidance on decontamination from the Microbiology Advisory Committee to Department of Health Medical Devices Agency". This document is more commonly known as the 'MAC Manual', and is referred to as such in this document for convenience.

With reference to the MAC Manual, "Introduction to Part 1 - Table 1: Classification of infection risk associated with the decontamination of medical devices"; many of the items of equipment comprising the Viamed Wall-Mounted Resuscitation Cabinet system are classified as low risk, due to being "In contact with healthy skin, or not in contact with the patient".

The recommended decontamination method for low risk items is to clean in accordance with the guidelines in the MAC Manual "Part 2; Cleaning (manual) - non-immersion".

Clean as per MAC Manual "Part 2; Cleaning (manual) – non-immersion"

- Radiant warmer
- Cabinet body
- Mounting rails
- Storage bins
- Integral bed
- Mattress
- Tom Thumb infant resuscitator
- Low suction controller
- VacSax suction receiving canister (can also be steam sterilized if required)
- Apgar timer

Cleaning Guidelines

The items listed above are low risk and not intended to be disinfected or sterilized, but where an alcohol wipe is used, this may have a disinfecting effect. After manual cleaning, an alcohol wipe containing 70% ethanol can be used over all external surfaces, observe safety precautions detailed later in this document.

It is recommended that a neutral detergent solution be used in accordance with the manufacturer's guidelines.

Sodium hypochlorite solutions are not recommended due to their potential corrosive effect on some metals.

The equipment within the cabinet system is mounted on standard mounting rails. To remove the equipment, loosen the rail clamps by rotating the locking nuts until the equipment can be lifted free of the rail.

It is recommended that the bed be cleaned in-situ, should it be necessary, the bed can be removed by removing the 6 screws holding it in place. Care should be taken when re-attaching the bed not to over-tighten the screws.

NOTE: the following instructions detail the method of manual cleaning (non-immersion) as directed by the MAC Manual. Should these instructions contradict guidelines in place within the customer's own organization, or those of the manufacturer of the cleaning detergent, please contact your Sterile Services Department or other responsible body within your organization for advice.

Equipment Required

- A warm water/detergent solution at correct dilution.
- A clean, disposable, absorbent, non-shedding cloth for application of detergent solution.
- A clean, disposable, absorbent, non-shedding cloth or mechanic drying facility (e.g. drying cabinet or industrial hot air dryer).
- An appropriate chemical neutraliser, first aid kit and eyewash bottle, in case of splashing with detergent.

Procedure

- If the item is electrical, ensure that it is disconnected from the mains supply before commencing the cleaning procedure.
- Wearing protective clothing, immerse the cleaning cloth in the detergent solution and wring thoroughly.
- Commencing with the upper surface of the item, wipe thoroughly ensuring that detergent solution does not enter electrical components.
- Periodically rinse the cloth in clean water and repeat the above steps.
- Surfaces should be carefully hand-dried using a cloth or industrial hot air dryer or placed into a drying cabinet.

Note: Non-immersion, manual cleaning is not a disinfection process, but where an alcohol wipe is used to dry surfaces, this may have a disinfecting effect.

- Safely dispose of cleaning materials and alcohol wipes, if used.

Monitoring and Control

Owing to the lack of control methods available to the user to test the efficiency of non-immersion cleaning, the user should be aware of the factors that may alter the efficiency of the method:

- staff training
- physical application
- nature of soil
- accessibility of cleaner to item/part of equipment
- detergent concentration

Safety Precautions

- Always wear protective waterproof clothing, robust gloves and eye protection if splashing is likely to occur.
- After removing protective clothing on completion of task, thoroughly wash and dry hands.
- Avoid splashing.
- Precautions should be taken when using alcohol, as it is flammable.
- The 'pooling' of alcohol on equipment should be avoided and alcohol evaporation ensured, if necessary by forced air drying. Care should also be taken to ensure that alcohol does not enter the item e.g. via ventilation slots.

Items that can be Steam Sterilized at 121° C as per HTM 2010, Part 3

- VacSax suction receiving canister
- VacSax suction taper
- Therapy Equipment reusable suction receiving jar

Items Intended for Single Patient Use Only

- Breathing circuit
- Suction tubing
- VacSax suction receiving liners
- Suction catheters



WARNING

Do not attempt to reprocess items intended for single patient use.

WARRANTY

All parts of the Viamed wall mounted infant resuscitation cabinet system are guaranteed for a period of 12 months from the date of purchase.

The best materials and workmanship have been employed throughout every stage of manufacture and every part is thoroughly tested before dispatch.

This warranty covers defects in material and manufacture but excludes damage caused by accident, misuse or neglect.

Should a component develop a defect within the warranty period, it will be repaired or replaced at Viamed's discretion.

In the event of warranty claims or queries regarding this product when purchased outside the UK, please contact your local distributor.

COMPANY DETAILS

All products are CE marked to the requirements of MDD 93/42 EEC and are supplied in accordance with our quality system accreditations:
BS EN ISO 9001/2000 and ISO 13485/2003.

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