Instructions for use

ASTOTHERM®*plus*

Warmer for Blood, Intravenous Fluids and Irrigation Fluids

REF AP220

Order No. 0550.7200.12 Rev. 06 10/2017

STIHLERELECTRONIC
STIHLER ELECTRONIC GmbH • 70597 Stuttgart • Germany

To be filled in by the user:

Serial number _____

Registration number _____

Device location _____

Start-up date _____

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CE 0124

STIHLER ELECTRONIC GmbH, Stuttgart, declares in sole responsibility that this product (230 – 240 VAC models only) conforms to EC Directive 93/42/EEC on medical devices.
Notified body: DEKRA Certification GmbH, registration number 0124.

Contents

1 Information about these Instructions

- **Carefully read the entire instructions for use before using the device.**
- **Correct and safe operation can only be guaranteed if the instructions for use are observed.**
- **Incorrect use can result in damage to the product or to other property and/or personal injury.**
- **Keep the instructions for use for future reference.**
- **Only use the device for the intended purpose as described in these instructions for use. Refer to *chapter***
- ***4 Specification of*** application***.***

2 General information

2.1 Guarantee conditions

The guarantee period is 12 months. During this guarantee period the manufacturer will repair or replace free of charge all defects caused as a result of material or manufacturing errors.

Other damage is not subject to this guarantee. The guarantee does not include cases of misuse or incorrect handling, use of force, or damage caused by normal wear and tear. This applies also to changes undertaken by persons who are not authorised by the manufacturer and to modifications to the original condition.

If the equipment is damaged during the guarantee period, send the cleaned equipment to the nearest sales point or directly to STIHLER ELECTRONIC GmbH. The sender is responsible for any transport and packaging costs.

2.2 Liability

The manufacturer is only liable for the safety, reliability and performance of the equipment

- if all operating, servicing, and calibration procedures have been carried out by trained and qualified persons according to the procedures published by the manufacturer;
- if only original spare parts have been used to replace components as needed;
- if assembly and repairs are only carried out by authorised personnel or an authorised service centre;
- if the electrical installations satisfy the locally applicable regulations and the IEC/EN requirements and
- if the equipment is used for its intended purpose and at a suitable location in accordance with the instructions for use.

2.3 Disposal of the equipment

Electrical devices are recoverable waste and should not be disposed of in domestic waste at the end of their service life. Please follow the local rules for the disposal of used products, or send the cleaned and disinfected equipment with a corresponding note to STIHLER ELECTRONIC GmbH or your closest sales point. This will ensure the most cost efficient and proper disposal of your old equipment.

	Follow the national regulations on the disposal of medical products.

2.4 Return of a used product

A report must be sent together with the equipment, detailing the precise reasons, circumstances, and, if known, the cause of the return. To prevent transportation damage, the equipment should be shipped either in the original packaging or in other, well-protected packaging.

 **WARNING**
Risk of infection! Clean and disinfect the equipment after every use and before you return the equipment for repairs.

NOTICE
The customer is responsible for the proper packaging and labelling of returns.

2.5 Service information

For service or technical support, please contact your local sales point or the following:

Terumo BCT Europe N.V
Ikaroslaan 41
1930 Zaventem
Belgium
Phone: +32.2.715.05.90
Fax: +32.2.721.07.70

Terumo BCT, Inc.
10811 W. Collins Avenue
Lakewood, Colorado 80215
USA
Phone: 303.231.4357
877.339.4228
Fax: 303.542.5215

www.terumobct.com

3 Important safety information

These Instructions for Use indicate and define the following safety information:

DANGER
DANGER indicates a hazardous situation which, if not avoided, will result in death or serious injury.

WARNING
WARNING indicates a hazardous situation which, if not avoided, could result in death or serious injury.

CAUTION
CAUTION indicates a hazardous situation which, if not avoided, may result in minor or moderate injury.

NOTICE
NOTICE indicates a property damage message.

3.1 Dangers

DANGER
Risk of explosion! Do not use the ASTOTHERM PLUS in an environment at risk of explosion or in the presence of flammable anesthetics.

3.2 Warnings

WARNING
Risk of injury! • Use of the ASTOTHERM PLUS must be carried out under the supervision of a physician. • Read and observe all instructions, stickers, and accompanying documentation enclosed with the medical device. Failure to observe the instructions, including warnings and safety information, can result in incorrect handling, patient injury, injury to users or medical personnel, damage to the device, or material damage. • Operate and service this equipment only in accordance with the procedures described in these instructions and with the applicable standards, rules, and guidelines. The manufacturer shall not be responsible for the safety of users or patients if any actions/procedures other than those published are carried out during operation, servicing, or recurrent tests.

WARNING

Risk of injury!

- This device may only be operated by appropriately trained and medically qualified healthcare professionals.
- The service personnel must be appropriately trained and qualified.
- Do not use the ASTOTHERM PLUS until the following error conditions have been remedied through appropriate corrective action:
  - Damaged or worn cables or plugs.
  - Damaged housing, damaged or loose control panel.
  - Device has been exposed to mechanical impact / exposed to severe shock or exposed to liquid.
  - Alarm without knowing the cause.
  - Damaged or missing markings/safety signs/warnings on the warmer.
- If the yellow “Alarm” LED and the acoustic alarm signal are not activated automatically, when the device turns on by pressing the “Standby” Button, remove the device from service immediately.
- In the event of an overheating alarm proceed as outlined below:
  1. Ensure that the ASTOTHERM PLUS safety system has deactivated the heating function and that the temperature is dropping below 43°C. If the temperature is not dropping, stop the treatment to prevent fluid from returning to the patient. Remove the applicable tubing immediately from the heat exchanger. Further evaluation should be carried out by qualified medical personnel such as a physician before blood in the line can be reinfused.
  2. Consider the possible reasons for the alarm. For further information see ***chapter 10 Alarms and troubleshooting***. If in doubt, do not continue using the warmer.
- The mains cable should not touch the patient and should not hinder the treating personnel.
- The ASTOTHERM PLUS does not contain any parts the user can repair. Therefore, do not attempt to repair the ASTOTHERM PLUS yourself. Contact your local sales point.
- Any repairs (such as, but not limited to, changing the power supply cord) to the equipment may only be carried out by persons authorized and qualified by the manufacturer.
- Modifications to the device are not permitted.

⚠WARNING

Risk of haemolysis!
Make sure that the infusion line is not kinked.

⚠WARNING

Risk of air embolism!

- When fluids are warmed up, it is possible that gas may evolve (bubble form).
- Be aware of the potential for air emboli when using a blood and fluid warmer.
- Therefore prime all filters, lines and disposable sets before starting a treatment.
- Make sure all connections of the complete fluid stream are fixed tightly to prevent fluid leakages and inadvertent infusion of air into the fluid stream.
- Do not warm infusions containing soluted gas (e.g. bicarbonate).
- Extreme care should always be taken to ensure that a bolus o fair does not pass to the patient.

⚠WARNING

Risk of infection!

- Use aseptic procedures.
- Clean and disinfect the warmer after every use and before you return the warmer for repairs.

⚠WARNING

Risk of electric shock!

- To avoid the risk of electric shock, this equipment must only be connected to a supply mains with protective earth.
- Do not use mains adapters that interrupt the earth conductor.
- Do not open the ASTOTHERM PLUS housing.
- If several pieces of equipment are combined or connected together (e.g., in multiple socket outlets), the total leakage currents must not exceed the allowable limits (refer to the respective national regulations). Observe the requirements as stipulated in IEC/EN 60601-1 regarding medical electrical systems.
- All electrical installations must conform to the applicable electrical standards and the specifications defined by the manufacturer.
- Before every use, check to make sure that the ASTOTHERM PLUS is undamaged.
- The mains plug must be removed from the socket to fully disconnect the ASTOTHERM PLUS from the mains.

WARNING

Risk of radio interference!

- Use of this equipment adjacent to or stacked with other equipment should be avoided because it could result in improper operation. If such use is necessary, this equipment and the other equipment should be observed to verify that they are operating normally.
- Use of accessories, other than those specified or provided by the manufacturer of this equipment could result in increased electromagnetic emissions or decreased electromagnetic immunity of this equipment and result in improper operation.
- Portable RF communications equipment (including peripherals such as antenna cables and external antennas) should be used no closer than 30 cm (12 inches) to any part of the ASTOTHERM PLUS, including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

3.3 Cautions

CAUTION

Risk of injury!

- Federal law (USA) restricts this device to sale by or on the order of a physician.
- When fixing the warmer to a mounting device, pay attention to the max. load to avoid tilting. Using normal infusion stands ASTOTHERM PLUS may be mounted at a height of up to 165 cm. If you use the robust IV pole ASTOSTAND, the device can be mounted up to 180 cm.
- Use only approved infusion sets / infusion extensions (e. g. ASTOTUBE, see ***chapter 15 Ordering information, accessories and consumables***).

CAUTION

Risk of hypothermia!

- When ASTOTHERM PLUS is used, the patient’s body temperature must be montitored at regular intervals.
- The specified heating performance will only achieved by inserting the infusion extension into the entire heat exchanger.
- The temperatur control of the ASTOTHERM PLUS controls and monitors the current temperature of the heat exchanger, but not the patient’s body temperature.
- If the ASTOTHERM PLUS cannot be started or if the patient's temperature balance is insufficient, consider the use of alternative warming methods in order to avoid or reduce hypothermia or to improve the patient's well-being.

⚠CAUTION

Risk of radio interference!

- The essential performance can be lost or degrated due to EM disturbances. As a result, there is the possibility of hypothermia of the patient.
- According to the standard IEC/EN 60601-1-2, medical electrical equipment requires special precautions regarding electromagnetic compatibility (EMC) and needs to be installed and put into service according the EMC information provided.
- This device/system may cause radio interference or may disrupt the operation of nearby devices. It may be necessary to take mitigation measures, such as re-orienting or relocating of ASTOTHERM PLUS or shielding the location.

3.4 Notices

NOTICE

- To avoid damage to the warmer:
  - Do not immerse the warmer in liquid.
  - Do not disinfect the warmer with these methods:
    - steam (autoclave)
    - hot air
    - thermo-chemical cleaning solutions.
  - Refer to the specific instructions for use of the disinfectants.
- The customer is responsible for the proper packaging and labelling of returns.

4 Specification of application

4.1 Intended use

ASTOTHERM PLUS is a warmer for Blood, Intravenous Fluids and Irrigation Fluids. The application areas include blood transfusions, intravenous fluids, dialysis, hemofiltration and apheresis.

4.2 Intended medical indication

The warming of medical fluids with ASTOTHERM PLUS supports the prevention and therapy of hypothermia.

4.3 Contraindications

There are no known contraindications for warming blood, intravenous fluids and irrigation fluids.

4.4 Possible adverse effects

There are no known side effects, which are causally attributed to the warming of blood, intravenous fluids or irrigation fluids.

4.5 Intended patient population

There are no restrictions for the intended patient group.

4.6 Intended user profile

The ASTOTHERM PLUS Warmer is to be operated only by medically qualified and trained healthcare professionals.

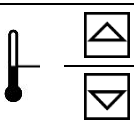
4.7 Intended use/operation environment

- The warmer may only be used in professional healthcare facilities (e.g. hospital, emergency care, dialysis, including HF surgical equipment, etc.).
- The warmer is not intended for home healthcare environment.
- The warmer is reusable, but requires cleaning / disinfection between the applications.
- Appropriate medical hygienic factors must be applied for the use of the warmer.
- The warmer must not be used in an environment at risk of explosion or in the present of flammable anesthetics.

4.8 Intended part of the body/type of tissue

The warmer is used to warm blood or other medical fluids supplied to the body. The fluids are physically separated from the warmer by disposable parts (tubes).

5 Symbols

Symbols, used on the Control Panel	
	Alarm condition if the yellow “Alarm” LED lights
	“Standby” button. The Warmer is in **Standby Mode** if the blue LED is on
	“Start” button. The Warmer is in **Heating Mode**, if the green LED is on
	“Increase” button increases the setpoint temperature
	“Decrease” button decreases the setpoint temperature

Where these symbols are applicable, they appear at the relevant point on the device, the package, the rating plate or in the accompanying paperwork.	
	Defibrillation-proof type B applied part in accordance with IEC/EN 60601-1
IPX 4	Splash proof in accordance to IEC/EN 60529
	Refer to instructions for use! / Follow instructions for use!
	Caution: Federal US law restricts this device to sale by or on order of a physician.
	General warning sign
	Catalogue number
	Serial number
	Year of manufacture
	Manufacturer
	Electrical devices are valuable products and should not be thrown in dustbin when the reach the end of their serviceable life.
	The device is conform to the MDD 93/42/EEC dated 14th June 1993 for medical devices. The Notified Body DEKRA Certifications GmbH (Reg. No. 0124) monitors the quality system of the manufacturer. The CE mark applies only to the ASTOTHERM PLUS Warmer. Disposable parts (e.g. infusion sets) suitable for use with this device have their own approvals.

CLASSIFIED c UL us 75JA	MEDICAL – GENERAL MEDICAL EQUIPMENT AS TO ELECTRICAL SHOCK, FIRE AND MECHANICAL HAZARDS ONLY IN ACCORDANCE WITH standards ANSI/AAMI ES60601-1:2005/(R)2012 and A1:2012), C1:2009/(R)2012 and A2:2010/(R):2012 CAN/CSA-C22.2 No. 60601-1:2014. Control No. 75JA
	This symbol indicates additional information.
	Indicates the temperature range within which the package must be stored and handled.
	Indicates the ambient humidity range within which the package must be stored and handled.
	Indicates the pressure range within the package must be stored and handled.
	Indicates the upright position of the package
	The package must be kept in a dry environment.
	The contents of the package are fragile and must therefore be handled with care. Do not drop or throw.
 06 PS	Recyclable - Polystyrene (acc. GB 18455-2001)
 CB	Recyclable - Cardboard (acc. GB 18455-2001)
	Acoustic alarm signal
	No acoustic alarm signal

6 Product description

6.1 Introduction

ASTOTHERM PLUS is a device for specific heating of blood and fluids which are delivered to the patients by transfusion, infusion or irrigation. The warming of the liquid supports the prevention and therapy of hypothermia during or after surgery; during longer procedures such as dialysis, hemofiltration, or apheresis the warming leads to a well-being. The applications of the ASTOTHERM PLUS therefore include transfusion, infusion, dialysis, hemofiltration and apheresis.

The ASTOTHERM PLUS Warmer can be used to warm fluids administered to a patient at flow rates of 0 to 6000 ml/h (i.e. 0 to 100 ml/min), see ***figure 1***.

The infusion extension ASTOTUBE is considered as applied part acc. to IEC/EN 60601-1.

6.2 Technical description

During operation of the Warmer heat is transferred from the internal heating element to the heat exchanger. Infusion extensions can simply be inserted in the circumferential groove. The heat from the heat exchanger is transferred through the inserted infusion extension to the fluid to be warmed.

The temperature of the heat exchanger is monitored by a microprocessor controlled temperature control system and by two independent alarm systems designed to alert the operator to failure conditions and, if necessary, to switch off the heating process automatically in the event of excessively high temperatures.

During operation, the mean temperature of the heat exchanger is displayed; this is not equal to the temperature of the medium to be warmed. ASTOTHERM PLUS neither regulates nor monitors/displays the current temperature of the medium to be warmed. The temperature of the medium (fluid) depends on a variety of factors, such as, but not limited to additional factors:

- Room temperature and ventilation
- Inlet temperature of the fluid (warmed-up or cold)
- Flow rate

CAUTION

Risk of hypothermia!

- When ASTOTHERM PLUS is used, the patient's body temperature must be montitored at regular intervals.
- The temperature control of the ASTOTHERM PLUS controls and monitors the current temperature of the heat exchanger, but not the patient's body temperature.
- If the ASTOTHERM PLUS cannot be started or if the patient's temperature balance is insufficient, consider the use of alternative warming methods in order to avoid or reduce hypothermia or to improve the patient's well-being.

Typical temperature curves are shown in the following figures.

10 ml/min = 0,6 l/h	60 ml/min = 3,6 l/h
20 ml/min = 1,2 l/h	70 ml/min = 4,2 l/h
30 ml/min = 1,8 l/h	80 ml/min = 4,8 l/h
40 ml/min = 2,4 l/h	90 ml/min = 5,4 l/h
50 ml/min = 3,0 l/h	100 ml/min = 6,0 l/h

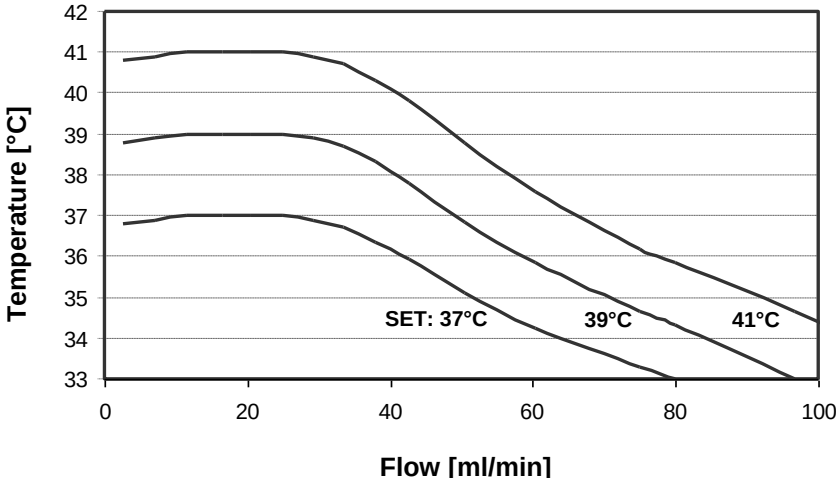


Fig. 1: Fluid outlet temperature at 20°C inlet temperature

6.3 Components of the ASTOTHERM PLUS

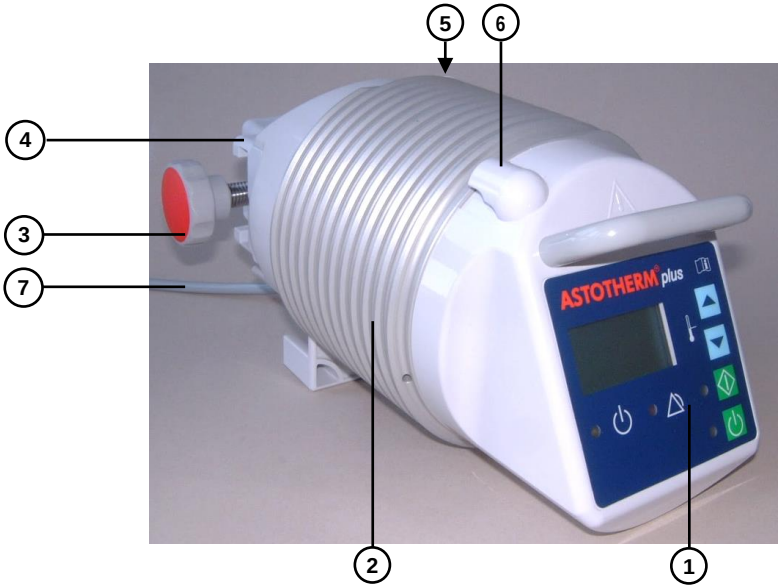


Fig. 2: ASTOTHERM PLUS AP220

#	Item	Description
1	**Control Panel**	Control buttons and displays. (***s. chapter 6.4 Control panel***)
2	**Heat exchanger**	Transfers heat from the internal heating element through the inserted infusion extension to the medium to be heated.
3	**Screw with Star Grip**	For adapting the attachment device to infusion stands of different diameters.
4	**Universal Attachment Device**	Attaches the warmer to infusion stands (Ø 12 to 35 mm) or medical rails.
5	**Tube holder rear**	Fixes the infusion extension at the entry point (from the liquid container).
6	**Tube holder front**	Fixes the infusion extension at the exit point (to the patient).
7	**Power Supply Cord with Mains Plug**	Conveys electricity from the wall power supply to the device. Pull the mains plug to disconnect from supply network.

6.4 Control panel

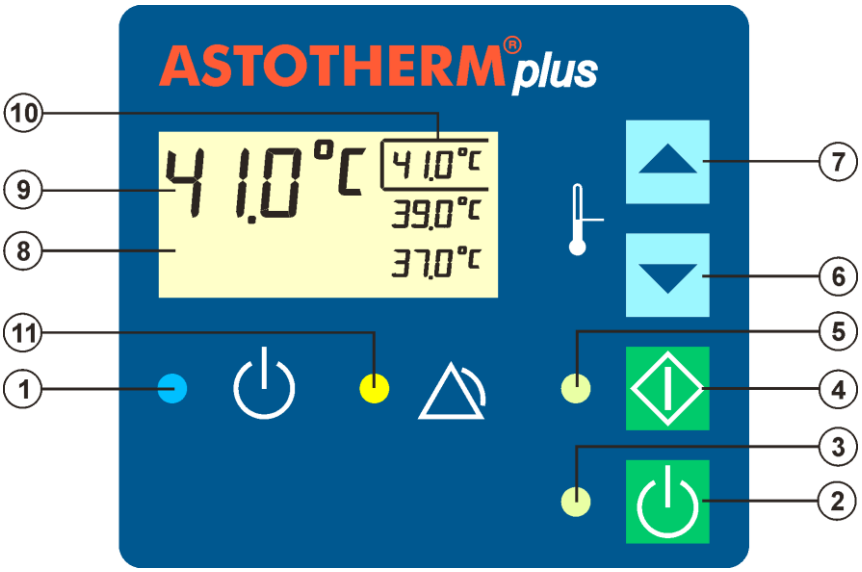


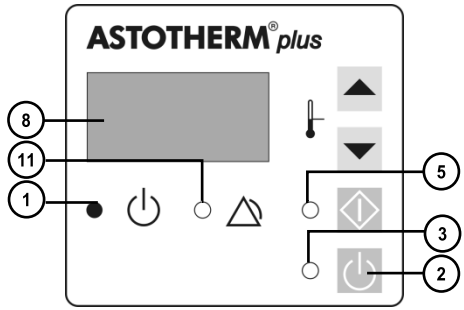
Fig. 3: Control panel

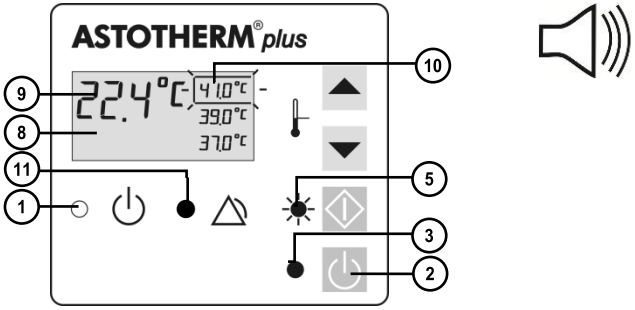
#	Item	Description
1	“Standby” LED	Illuminates blue when the warmer is in **Standby Mode**.
2	“Standby” button”	Turns the warmer from **Standby Mode** to **On Mode**.
		Turns the warmer from any **Mode** to the **Standby Mode**.
3	“On” LED	Illuminates green when the warmer is in **On Mode**.
4	“Start” button	Starts the heating process while the device is in **On Mode** or **Alarm Mode**.
		Starts test 6 (see ***chapter 12.1 Recurrent tests***), when the warmer is operated with the middle setpoint temperature.
5	“Start” LED	Flashes green when the device is in **On Mode** (heating is not yet started).
		Illuminates green when the device is in **Heating Mode** (“Start” button has been pressed).
6	“Decrease” button	Selection of the next lower setpoint temperature. The frame indicates the selected temperature.
		Starts test 8 (see ***chapter 12.1 Recurrent tests***), when the warmer is operated with the lowest setpoint temperature.
7	“Increase” button	Selection of the next higher setpoint temperature. The frame indicates the selected temperature.
		Starts test 7 (see ***chapter 12.1 Recurrent tests***), when the warmer is operated with the highest setpoint temperature.

#	Item	Description
8	LCD Display	Informs the user about temperatures, test and fault conditions.
9	Actual Temperature	Displays the current temperature of the heat exchanger.
10	Setpoint Temperatures	Shows the three possible setpoint temperatures. The frame indicates the selected temperature.
11	“Alarm“ LED	Lights yellow if an alarm condition exists.

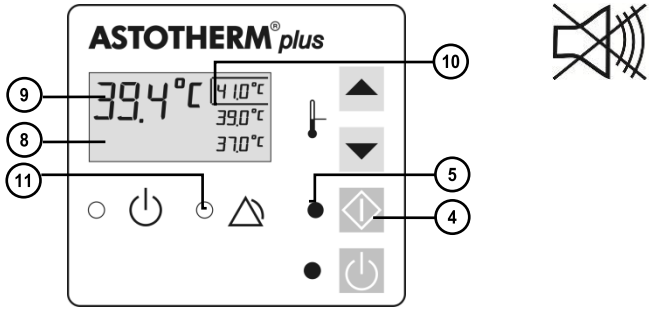
The following section provides further information about the operating states. This includes a description of the actions of the user and the device responses of each operating state.

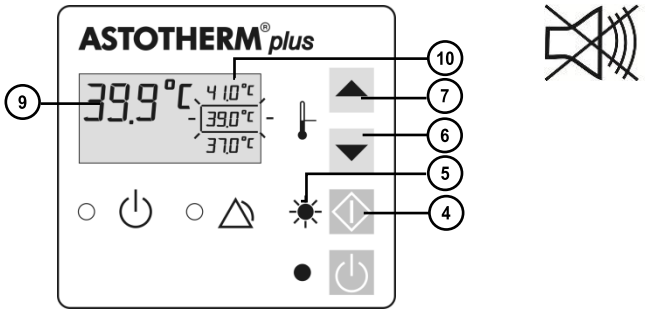
7 Operating states

7.1 Standby mode	
Control Panel	
Action	When the mains cable is plugged into the socket, the device is in **Standby Mode**. Alternately, press the “Standby” button ② to switch the device from any **Mode** to **Standby Mode**.
Device response	When plugging in the power cord briefly all segments of the display and all LEDs turn on, then  • all segments of the display turn off ⑧. • “Start” LED ⑤, “On” LED ③, “Alarm” LED ⑪ turn off. • “Standby” LED ① turns on.
ⓘ	• After a prolonged power failure (> 5 sec), the device automatically switches to **Standby Mode**. • In **Standby Mode**, only the electronic is disconnected from the power supply. The Warmer remains connected to the mains.

7.2 On mode	
Control Panel	
Action	Press the “Standby” button (2) to switch the device from **Standby Mode** to **On Mode**.
Device response	• “Standby” LED (1) turns off. • “On” LED (3) turns on. • The backlight of the display (8) lights up. • The display (8) shows the current temperature (9) (eg. 22.4°C) of the heat exchanger and the available range of setpoint temperatures (10). The selection frame is flashing. • “Start” LED (5) flashes. • „Alarm” LED (11) lights • Acoustic alarm signal beeps.
	As long as the temperature of the heat exchanger is below 15°C, the display shows “- - -”.

7.3 Heating mode

Control Panel	
Action	Press the “Start” button (4) at least for one second to switch the warmer from **On Mode** to **Heating Mode**.
Device response	• While the button is pressed, a self-test is performed. During this test the safety cut offs are activated to verify their safe function. The brief clicking of the relay can be heard. • “Start” LED (5) turns on. • Acoustic alarm signal stops. • The heater is activated until the setpoint temperature (10) (indicated by the frame) is reached. • The display (8) shows the increasing current temperature (9) of the heat exchanger (e.g. 39.4°C). • “Alarm” LED (11) lights until the setpoint temperature has exceeded (during heating up) the tripping limit of the Low Temperature Alarm (see ***chapter 10.1 Low temperature alarm***).
	• If the “Start” button“ (4) is not pressed long enough, the self-test cannot be completed and the warmer will not start. Then repeat the procedure and press the “Start” button (4) for at least one second. • After a short power failure (< 5 seconds), and after return of the power supply, the warmer resumes operation again to the setpoint temperature which was previously selected.

7.4 Increasing/Decreasing the setpoint temperature	
Control Panel	
Action	1. Briefly press the “Increase” button ⑦ or “Decrease” button ⑥, while the device is turned on (**On Mode**) or started (**Heating Mode**) to change the setpoint temperature within the three indicated temperatures. 2. Confirm the new setpoint temperature within 5 seconds by pressing the “Start” button ④.
Device response	• The flashing frame indicates the selected setpoint temperature ⑩. • “Start” LED ⑤ flashes green until to the confirmation with the “Start” button ④. • After confirmation the “Start” LED ⑤ lights green and the frame marks the currently selected setpoint temperature ⑩. • The displayed actual temperature ⑨ rises or falls according to the selection.
ⓘ	• If the newly selected setpoint temperature is not confirmed within 5 seconds by pressing the “Start” button ④, the temperature regulation continues working with the initial setpoint temperature. • During operation, the mean temperature of the heat exchanger is displayed; this is not equal to the temperature of the medium to be warmed. ASTOTHERM PLUS neither regulates nor monitors/displays the current temperature of the medium to be warmed.

8 Installation

8.1 Initial start-up

Prior to first use, perfom the following inspections:

- Visual inspection (s. ***chapter 12.1 Recurrent tests***)
- Check the mains voltage (compare the details on the type label with the available mains voltage.) An incorrect mains voltage may destroy the equipment.

National regulations may require different inspections for the initial start-up. If additional tests are required for electrical safety, these must be carried out according to ***chapter 12.1 Recurrent tests, 12.2 Set up for electrical tests and 12.3 Test protocol***.

8.2 Installation of the warmer

For safe installation, the warmer is equipped with a universal attachment device. With this, the device can be securely attached to infusion stands as well as to medical standard rails.

8.2.1 Attachment to infusion stands/rods

1. Turn the handwheel counterclockwise to open the attachment device.
2. Select a maximum height of 165 cm (ASTOSTAND: 180 cm) at the infusion stand and place the open clamping area of the attachment device on the infusion stand.
3. Turn the handwheel clockwise to lock the attachment device to the infusion stand.
4. Check that the warmer is firmly fixed.

8.2.2 Attachment to medical rails

1. Unscrew the small knurled screw on the bottom of the attachment device.
2. Hang the warmer obliquely from above with the attachment device into the standard rail.
3. Fix the heater by tightening the small knurled screw to the standard rail.
4. Check that the warmer is firmly fixed.

9 Getting started

Getting started is grouped into 4 sections. Read through each section before performing a procedure.

CAUTION

Risk of injury!
Use only approved infusion sets / infusion extensions.

ASTOTUBE is the original CE-marked accessory for ASTOTHERM PLUS.

ASTOTUBE Order No.	Description	Suitable for model
IFT 40410	Sterile infusion extension made from PVC, outer diameter Ø 4 mm, length 592 cm. Filling volume approx. 42 ml	**AP220**

Do not position the device in a manner that is difficult to disconnect it from mains by the mains plug.

9.1 Preparation for use

WARNING

Risk of injury!

- Do not use the ASTOTHERM PLUS until the following error conditions have been remedied through appropriate corrective action:
  - Damaged or worn cables or plugs.
  - Damaged housing, damaged or loose control panel.
  - Device has been exposed to mechanical impact / exposed to severe shock or exposed to liquid.
  - Alarm without knowing the cause.
  - Damaged or missing markings/safety signs/warnings on the warmer.
- Use of the ASTOTHERM PLUS must be carried out under the supervision of a physician.
- The mains cable should not touch the patient and should not hinder the treating personnel.

CAUTION

Risk of injury!
When fixing the Warmer to a mounting device (e.g. IV pole) pay attention to the max. safe working load and tipping stability. Using normal infusion stands ASTOTHERM PLUS may be mounted at a height of up to165 cm. If you use the robust IV pole ASTOSTAND, the device can be mounted up to 180 cm.

1. Attach ASTOTHERM PLUS to an IV pole or to a medical standard rail using the attachment device according to ***chapter 8.2 Installation of the warmer***.
2. Plug the mains plug into a socket.
   - The blue “Standby” LED turns on, the device is in **Standby Mode**.
3. Press the “Standby” button to switch ASTOTHERM PLUS to **On Mode**.
   - The blue “Standby” LED turns off and the green “On” LED turns on.
4. Check the audible and visual indicators and the display:
   - The acoustic alarm signal beeps and the “Alarm” LED lights yellow.
   - The „Start” LED flashes green and the display shows the actual temperature of the heat exchanger and the available setpoint temperatures.

WARNING
Risk of injury! If the yellow "Alarm” LED and the acoustic alarm are not activated automatically when the device turns on by pressing the “Standby” button, remove the device from service immediately.

5. Press the “Increase” button or the “Decrease” button to select a different setpoint temperature, if necessary.
6. Press the “Start” button for at least one second to switch ASTOTHERM PLUS to **Heating Mode**.
   - While the button is pressed, the clicking of the self-test can be heard.
   - The “Start” LED illuminates green.

	• Any change of the setpoint temperature must be confirmed within 5 seconds by pressing the „Start” button, otherwise the warmer uses the previously selected setpoint temperature. • As long as the temperature of the heat exchanger is below 15°C, the display shows “- - -“. • As long as the actual temperature is below the trigger temperature of the low temperature alarm (4°C lower than the selected setpoint temperature), the “Alarm” LED lights yellow. • The setpoint temperature can be changed during operation at any time following ***chapter 7.4 Increasing/Decreasing the setpoint temperature***.

9.2 Inserting the infusion line, priming and starting the infusion

WARNING

Risk of air embolism!

- Be aware of the potential for air emboli when using a blood and fluid warmer.
- Therefore prime all filters, lines and disposable sets before starting a treatment.
- Make sure all connections of the complete fluid stream are fixed tightly to prevent fluid leakages and inadvertent infusion of air into the fluid stream.
- When fluids are warmed up, it is possible that gas may evolve (bubble form).
- Do not warm infusions containing soluted gas (e.g. bicarbonate).
- Extreme care should always be taken to ensure that a bolus of air does not pass to the patient.

WARNING

Risk of infection!

Use aseptic procedures.

WARNING

Risk of haemolysis!

Make sure that the infusion line is not kinked.

CAUTION

Risk of hypothermia!

- When ASTOTHERM PLUS is used, the patient’s body temperature must be montitored at regular intervals.
- The specified heating performance will only achieved by inserting the infusion extension into the entire heat exchanger.
- The temperatur control of the ASTOTHERM PLUS controls and monitors the current temperature of the heat exchanger, but not the patient’s body temperature.
- If the ASTOTHERM PLUS cannot be started or if the patient's temperature balance is insufficient, consider the use of alternative warming methods in order to avoid or reduce hypothermia or to improve the patient's well-being.

Pressure losses can occur when infusion extensions are used (depending on tube dimensions and flow rate).

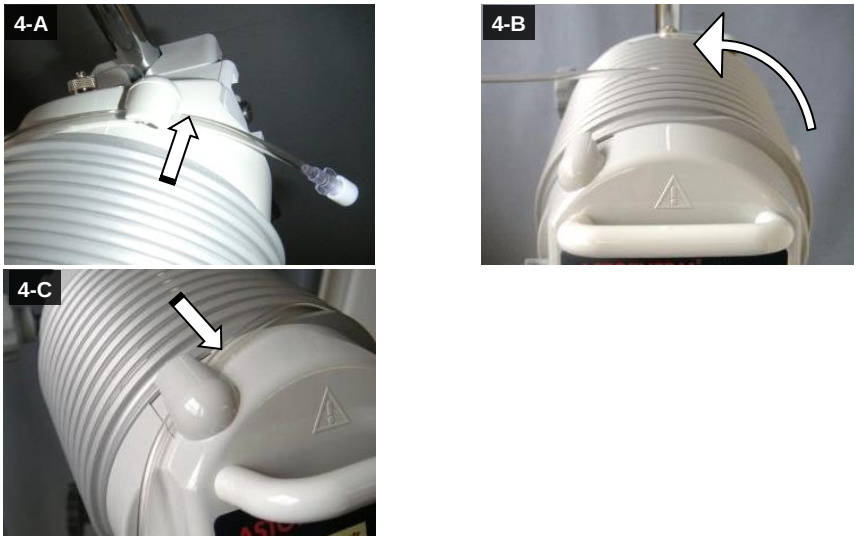


Fig. 4: Inserting the infusion extension

1. Clip the IV end of the infusion extension (female Luer Lock) into the rear tube holder (fig. 4-A).
2. Starting from the back, wind and insert the infusion extension in a counter clockwise direction up and around into the circumferential groove of the heat exchanger (fig. 4-B). Gently pulling makes it easier to insert and improves the seating of the infusion extension.
3. After leaving the last circulation, clip the infusion extension into the front tube holder (fig. 4-C).
4. Check the correct position of the infusion extension:
   a. Infusion extension is fully in the groove
   b. Infusion extension „skips“ no circulation
   c. Infusion extension is not kinked or twisted in itself
5. Connect the infusion extension with infusion set of the liquid container.
6. Prime the infusion system: allow fluid to flow until no air is observed in the infusion lines and the lines are completely filled with fluid.
7. Connect the patient side end of the infusion extension to the patient cannula and ensure a good fixation (e.g. with tape).

Fig. 5: ASTOTHERM PLUS prepared

The infusion extension is now free hanging between the patient and ASTOTHERM PLUS and the treatment can be started.

9.3 After use

1. Discontinue infusion.
2. Press the “Standby” button for one second to switch off ASTOTHERM PLUS.
   - All indicators turn off, the LED “Standby” turns on blue.

	To disconnect ASTOTHERM PLUS from the mains, it is necessary to completely pull out the plug.

3. Disconnect the infusion extension from the cannula.
4. Wrap the infusion extension out from the circumferential groove of the heat exchanger.
5. Clean and disinfect ASTOTHERM PLUS after each treatment and if required.

9.4 Cleaning and disinfecting

NOTICE
To avoid damage to the warmer: - Do not immerse the warmer in liquid. - Do not disinfect the warmer with these methods: ▪ steam (autoclave) ▪ hot air ▪ thermo-chemical cleaning solutions. - Refer to the specific instructions for use of the disinfectants.

Clean and wipe-disinfect ASTOTHERM PLUS in accordance with the procedure below:

1. Disconnect the mains plug from the socket.
2. Clean all surfaces with a soft cloth/cotton swab and mild soap-and-water solution.
3. Disinfect **ASTOTHERM PLUS** with either:
   - An approved disinfectant
   - An alcohol-based disinfectant with a low content of aldehydes (< 0.2%)
   - A mild bleach solution (max. 0.25 % hypochlorite)

List of approved disinfectants*:

• Meliseptol®	• Clinell Alcohol Wipes
• Biguamed® Perfekt N	• Incidin® Plus
• Mikrozid® Liquid	• HyPro medical 3% H_2O_2
• Bacillol® Plus	• Aniosurf
• Mikrobac® forte	• Oxivir Tb
• ClearSurf®	• Diosol 3% H_2O_2 PURE
• Clinell Universal Sanitising Wipes	• Virox5 RTU

*In the United States please use only disinfectants which are registered by EPA (U.S. Environmental Protection Agency) or cleared by FDA (U.S. Food and Drug Administration).

10 Alarms and troubleshooting

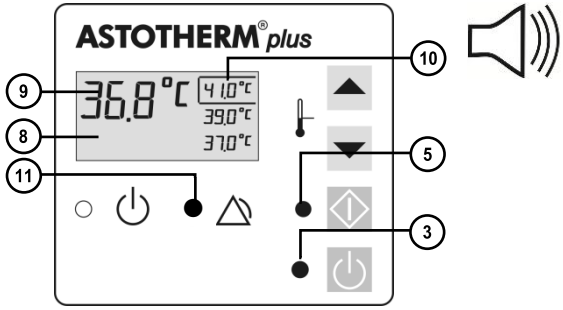
In the event of device failure, two independent monitoring systems ensure the safety against overheating. Except for the low temperature alarm, all alarms switch off immediately the heating function. Thus, the overheating of the heated liquid is surely prevented.

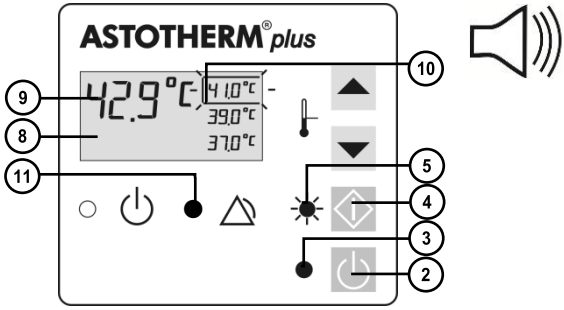
ASTOTHERM PLUS does not require continuous supervision by the operator, but it must be checked at regular intervals (depending on the condition of the patient status). Then the intended operator's position is directly in front of the control panel.
In case of failure of the equipment, possible injury to the patient is delayed and the operator has sufficient time to provide alternative warming methods.

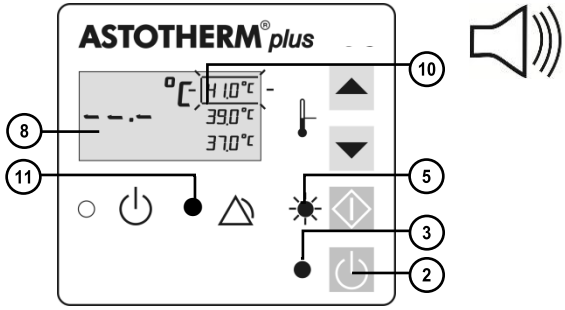
According to IEC/EN 60601-1-8 the alarms are defined as "**Low Priority Alarms**".

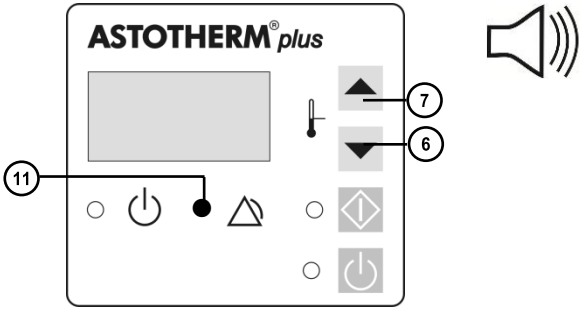
The alarms are only triggered by technical alarm conditions (equipment faults). The alarm signal is given visually and acoustically.

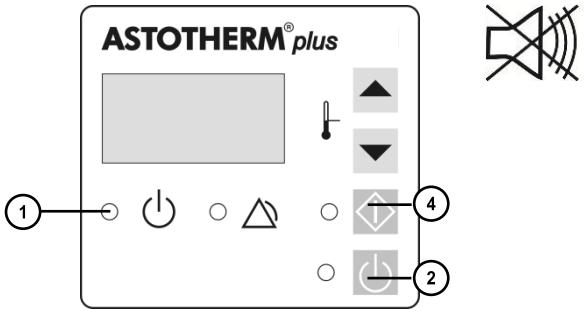
Alarm signal	**Characteristics**
visible	yellow LED lights constantently
audible	Sound pulse, all 17 sec.

10.1 Low temperature alarm	
Control panel	
Device response	• The display (8) shows an actual temperature (9), which is more than 4°C below the setpoint temperature (10). • The green “On” LED (3) lights. • The green “Start” LED (5) lights. • The “Alarm” LED (11) lights yellow. • The acoustic alarm signal is activated with a 2 minute delay. • The heating element is not switched off.
Alarm condition	The current temperature of the heat exchanger is more than 4°C below the setpoint temperature during **Heating Mode**.
Possible reason(s) ►Required action(s)	Inlet temperature of the liquid is too low and the flow rate is too high. ►Reduce flow rate.
	Warmer is defective. ►Return ASTOTHERM PLUS to local sales office.
Possible actions to clear	None, the alarm is automatically disabled if the alarm condition is eliminated.
ⓘ	During warm up phase, as long as the actual temperature is 4°C below the selected setpoint temperature, low temperature alarm is displayed (visual alarm signal only).

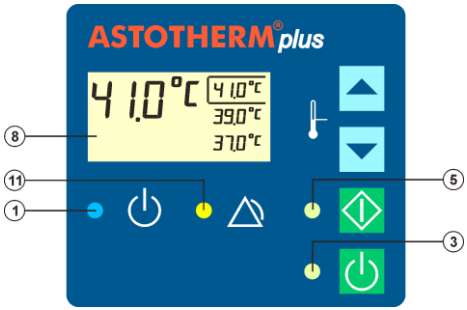
10.2 Overheating alarm	
Control panel	
Device response	• The display ⑧ shows the current temperature ⑨, which drops slowly after the alarm was activated. • The selection frame ⑩ flashes. • The green “On” LED ③ lights. • The “Start” LED ⑤ flashes green. • The “Alarm” LED ⑪ lights yellow. • The acoustic alarm signal sounds every 17 seconds. • The heating element is switched off.
Alarm condition	The temperature of the heat exchanger rises to the alarm limit of **45.5°C ± 1.0°C**.
Possible reason(s) ►Required action(s)	Effect of external heat source, such as sunlight or radiator. ►Eliminate the heat source and/or select cooler location.
	Ambient temperature too high. ►Eliminate heat source and/or select cooler location.
	Warmer is defective. ►Return ASTOTHERM PLUS to local sales office.
Possible actions to clear	Press the “Start” button ④ to switch the device back to **Heating Mode** (after cooling down).
	Press the “Standby” button ② to switch the device to **Standby Mode**.
	• As long as the actual temperature ⑨ is above the alarm limit, the warmer cannot be switched to the **Heating Mode**. • In order to prevent potential overheating due to a possible failure of the temperature control system, the ASTOTHERM PLUS possesses two independent excessive temperature cut offs.

10.3 Cable break alarm	
Control panel	
Device response	• The display (8) shows „- - -„. • The selection frame (10) flashes. • The green “On” LED (3) lights. • The “Start” LED (5) flashes green. • The “Alarm” LED (11) lights yellow. • The acoustic alarm signal sounds every 17 seconds. • The heating element is switched off.
Alarm condition	A defect of the cable break detection or a temperature sensor break has been detected.
Possible reason(s) ►Required action(s)	Warmer is defective. ►Return ASTOTHERM PLUS to local sales office.
Possible actions to clear	Press the “Standby” button (2) to switch the device to **Standby Mode**.

10.4 Processor alarm	
Control panel	
Device response	Even when you plug in the power cord …. • the “Alarm” LED (11) lights yellow. • the acoustic alarm signal sounds every 17 seconds. • none of the buttons can cause a device reaction.
Alarm condition	Program fault.
Possible reason(s) ►Required action(s)	Temporary program fault. ►Alarm reset (see below).
	Permanent program fault caused by defect data file. ►Return ASTOTHERM PLUS to local sales office.
Possible actions to clear	1. Press the “Increase” button (7) and the “Decrease” button (6) at the same time until the device switches to **Standby Mode**. 2. Unplug the power cord and wait one minute. 3. Plug in again the power cord.

10.5 Standby mode - failure	
Control panel	
Device response	The “Standby” LED ① is off and the device cannot be switched to **On Mode** by pressing the “Standby” button ②.
Possible reason(s) ►Required action(s)	Power supply problem or no power. ►Check plugs and fuses compare mains voltage with rating plate.
	Warmer not plugged in. ► 1. Plug the warmer into functioning socket. 2. Press the “Standby” button ② . 3. Press then the “Start” button ④ to switch the device to **Heating Mode**.
	Warmer is defective. ►Return ASTOTHERM PLUS to local sales office.

11 Brief overview of operating states and alarms



Operating state	Display (8)	"Standby" LED blue (1)	"Alarm" LED yellow (11)	"Start" LED green (5)	"On" LED green (3)	Acoustic alarm signal	Possible reason(s)
Standby Mode	*OFF*	●	○	○	○	🔇	-
On Mode	---	○	●	☀	●	🔊	$T_{Act} \leq 15°C$ or $T_{Act} \geq 50°C$
	T_{Act}	○	●	☀	●	🔊	-
Heating Mode	---	○	●	●	●	🔇	$T_{Act} \leq 15°C$
	T_{Act}	○	○	●	●	🔇	-
	T_{Act}	○	●	☀	●	🔊	„Start" button has been pressed too short. Otherwise device defect.

T_{act} = Actual Temperature (current temperature of heat exchanger)
T_{set} = Setpoint Temperature (selected temperature, marked by a frame)

○ = LED is off ● = LED lights ☀ = LED flashes

Alarm	Display (8)	"Standby" LED blue (1)	"Alarm" LED yellow (11)	"Start" LED green (5)	"On" LED green (3)	Acoustic alarm signal	Possible reason(s)
Low Temperature Alarm	T_{act}	○	●	●	●	sounds every 2 min.	Low temperature ($T_{act} \leq T_{set}$ - 4°C) because of cold liquid/high flow rate or device defective
Excessive Temperature Alarm	T_{act}	○	●	☀	●	🔊	T_{act} > 45.5°C ± 1°C
Cable Break Alarm	---	○	●	☀	●	🔊	Temperature sensor(s) or associated circuit(s) interrupted
Processor Alarm	OFF	○	●	○	○	🔊	Program failure
Manual Excessive Temperature Alarm Test 1	t 1 alternating with T_{act}	○	●	☀	●	🔊	„Start" button has been pressed for longer than 3 seconds
Manual Excessive Temperature Alarm Test 2	t 2 alternating with T_{act}	○	●	☀	●	🔊	„Increase" button has been pressed for longer than 3 seconds
Manual Low Temperature Alarm Test 3	t 3 Alternating with T_{act}	○	●	●	●	🔊 after cooling to $T_{act} \leq T_{set} - 4°C$	„Decrease" button has been pressed for longer than 3 seconds

T_{act} = Actual Temperature (current temperature of heat exchanger)
T_{set} = Setpoint Temperature (selected temperature, marked by a frame)

○ = LED is off ● = LED lights ☀ = LED flashes

12 Maintenance

ASTOTHERM PLUS does not require preventive maintenance (e.g. replacement of liquids or components). Recurrent tests shall be performed according to chapter 12.1.

ⓘ	No service or maintenance shall be carried out while in use with a patient.

⚠WARNING

- **Risk of injury!**
- The service personnel must be appropriately trained and qualified.
- The ASTOTHERM PLUS does not contain any parts the user can repair. Therefore, do not attempt to repair the ASTOTHERM PLUS yourself. Contact your local sales point.
- Any repairs (such as, but not limited to, changing the power supply cord) to the equipment may only be carried out by persons authorised and qualified by the manufacturer.
- Modifications to the device are not permitted.

The accessories specified in ***chapter 15 Ordering information, accessories and consumables*** may be replaced without restriction by the operating or maintenance personnel.

On request, STIHLER ELECTRONIC GmbH will provide service instructions that will allow properly trained and qualified persons to repair the parts of the equipment that the manufacturer has designated as repairable.
Provision of technical documents and/or spare parts is not an authorisation from the manufacturer to open or repair the equipment.

12.1 Recurrent tests

A recurrent test must be carried out on the ASTOTHERM PLUS warmer at least every 24 months to ensure the safe operation of the warmer.
Please ensure that all the applicable national directives (e.g. IEC/EN 62353) for checking the safety of medical equipment are observed additionally and that the test equipment is calibrated.

Necessary test equipment:
- Standard medical electrical safety tester
- Digital clinical thermometer (max. diameter of probe tip 3.5 mm and accuracy ± 0.1°C)

The following sections describe how the tests are to be performed. The attached test protocol (see ***chapter 12.3 Test protocol***) can be used.

Test 1	**Visual inspection**
Required Actions	Check the following items: • Complete and legible labeling. • No damage to the housing. • Control panel in good condition. (Since the front foil prevents fluid from entering the unit, it is important that its full surface adheres securely to the housing.) • No defects in the power supply cord and mains connector insulation, with clean and non-corroded contacts.

Test 2	**Protective earth resistance**
Required Actions	Measure the resistance between the earth pin on the power plug and the connector for potential equalization, located on the rear side of the housing. For detailed information performing this test see ***chapter 12.2 Set up for electrical tests***.
Result	The test is successful when the limits are met in accordance with the test protocol.

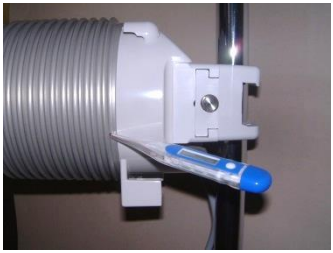
Test 3	**Insulation resistance**
Required Actions	Measure the resistance of the insulation between the live parts and the parts which are connected to earth. For detailed information performing this test see ***chapter 12.2 Set up for electrical tests***.
Result	The test is successful when the limits are met in accordance with the test protocol.

Test 4.1 Optional to test 4.2	**Equipment leakage current (alternative method)**
Required Actions	Measure the current flowing from protective earthed conductor to the two (shorted) power supply connections. For detailed information performing this test see ***chapter 12.2 Set up for electrical tests***.
Result	The test is successful when the limits are met in accordance with the test protocol.

or alternatively:

Test 4.2 Optional to test 4.1	**Earth leakage current (direct method)**
Required Actions	Measure the maximum earth leakage current (PE open)*. Measure all combinations of line polarity with neutral open (single fault condition) and closed (normal condition). For detailed information performing this test see ***chapter 12.2 Set up for electrical tests***.
Result	The test is successful when the limits are met in accordance with the test protocol.

*is usually done automatically by the safety tester

Test 5	**Temperature control and display**
Required Actions	1. Insert the clinical thermometer into the rear measuring bore on the side of the heat exchanger.  **Fig. 6: Temperature measurement** 2. Plug the mains plug into a socket. 3. Press the “Standby” button (**On Mode**). 4. Select a temperature of maximum 41°C (if necessary press the “Decrease” button or the „Increase” button . 5. Press the “Start” button for at least one second (**Heating Mode**). 6. Wait about 5 minutes until the actual temperature is equal to the setpoint temperature. 7. Start the measurement on the clinical thermometer and measure the actual temperature of the heat exchanger. 8. Compare the measured temperature with the displayed temperature and the selected setpoint temperature of the Warmer.
Result	The test is successful when the limits are met in accordance with the test protocol (***chapter 12.3 Test protocol***)
	• This test is used to check the essential performance. • Absolutely avoid during this measurement influences from the environment (drafts, heat radiation of other heat sources, etc). • Clinical thermometers are designed as „immersion sensors“. In order to achieve a sufficiently accurate measurement result, the thermometer must be immersed deep enough (depending on manufacturer and type). Because here only the metal tip of the thermometer is used, usually the measured temperature is slightly lower than the real temperature.

Test 6	**Manual excessive temperature cut off 1**
Required Actions	1. Operate the Warmer with the middle setpoint temperature. 2. Hold down the “Start” button at least for 3 seconds, to start the test.
Result	The test is successful, when: • The display alternately indicates **t1** and the actual temperature and after a short time – the “Start” LED flashes green, – the “Alarm” LED lights yellow, – the acoustic alarm signal sounds. The test is not successful if any of the following conditions occurs: • The display does not indicate **t1**. • The “Start” LED does not flash. • The “Alarm” LED does not light. • The acoustic alarm does not beep.
	For the continuation of the tests, press the “Start” button to switch the Warmer back to the **Heating Mode**.

Test 7	**Manual excessive temperature cut off 2**
Required Actions	1. Operate the Warmer with the highest setpoint temperature. 2. Hold down the “Increase” button at least for 3 seconds, to start the test.
Result	The test is successful, when: • The display alternately indicates **t2** and the actual temperature and after a short time – the “Start” LED flashes green, – the “Alarm” LED lights yellow, – the acoustic alarm signal sounds. The test is not successful if any of the following conditions occurs: • The display does not indicate **t2**. • The “Start” LED does not flash. • The “Alarm” LED does not light. • The acoustic alarm does not beep.
	For the continuation of the tests, press the “Start” button to switch the Warmer back to the **Heating Mode**.

Test 8	**Manual low temperature alarm**
Required Actions	1. Operate the Warmer with the lowest setpoint temperature. 2. Hold down the “Decrease” button ▽ at least for 3 seconds, to start the test.
Result	The test is successful, when: • The display alternately indicates **t3** and the actual temperature. • The actual temperature slowly decreases, and after cooling to $T_{Act} = T_{Set} – 4°C$ – the “Alarm” LED △ flashes yellow and – after another 2 minutes, the acoustic alarm signal sounds. The test is <u>not</u> successful if any of the following conditions occurs: • The display does not indicate **t3**. • The “Alarm” LED △ does not light after cooling. • The acoustic alarm signal does not beep after another 2 minutes waiting time.
ⓘ	• The cooling time depends in the initial temperature and the ambient temperature. • To exit the test, press the “Standby” button ⏻ to switch the Warmer to the **Standby Mode**.

12.2 Set up for electrical tests

For measuring the protective earth resistance, the insulation resistance and the equipment/earth leakage current, the following test set up can be used:

Test	Measurement (True RMS) (see also IEC/EN 62353)	Use connection to electrical safety analyzer
2	Protective earth resistance	Connection 1 and 2
3	Insulation resistance	Connection 1
4.1 *optional to 4.2*	Equipment leakage current (alternative method)	Connection 1 (and possibly connection 2, depending on the used safety analyzer)
4.2 *optional to 4.1*	Earth leakage current N.C. (direct method)	Connection 1
	Earth leakage current S.F.C (direct method, N open)	

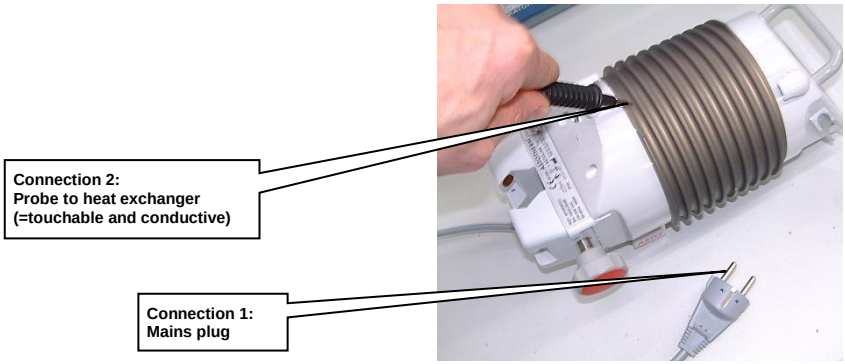


Abb. 7: Test set up for measuring

ⓘ	In order to achieve a sufficient accurate measurement result, during the measurement of the protective earth resistance, good electrical contact with the metal of the heat exchanger must be established. On the top of the heat exchanger (backside) is a small bare spot to make the contact with the probe. At this point the (insulating) anodic coating of the heat exchanger is penetrated by the manufacturer's test.

12.3 Test protocol

	ASTOTHERM PLUS
Type	
SN	

Test equipment		
Type		
SN		
Date of calibration		

Test 1: Visual inspection	P/F
Labeling and markings on ASTOTHERM PLUS	
Control panel (front foil)	
Housing	
Power supply cord	
Attachment device	

Test 2: Protective earth resistance			
	Value [Ω]	**Max [Ω]**	**P/F**
Protective earth resistance		0.3	

Test 3: Insulation resistance			
	Value [MΩ]	**Min [MΩ]**	**P/F**
Insulation resistance		100	

Test 4: Leakage current test either to 4.1 or 4.2			
☐ **4.1 Equipment leakage current (alternative method)** Alternative test for measurement the earth leakage current using the direct method (Test 4.2)			
	Value [mA]	**Max [mA]**	**P/F**
Equipment leakage current		1.0	
☐ **4.2 Earth leakage current (direct method)** Alternative test for measurement the equipment leakage current using the alternative method (Test 4.1) PE (Protective ground) open. Measure all combinations of line polarity.			
	Value [mA]	**Max [mA]**	**P/F**
Earth leakage current N.C		0.5	
Earth leakage current S.F.C (N open)		1.0	

Test 5: Temperature control and display				
	Value [°C]	**Min [°C]**	**Max [°C]**	**P/F**
Selected setpoint temperature **T** (max. 41°C)				
Temperature measurement **TT** (with clinical thermometer)		**T** – 0.5	**T** + 0.5	
Actual temperature (displayed) **TD**		**T** – 0.3	**T** + 0.3	

Manual tests	**P/F**
Test 6: Manual excessive temperature cut off 1 (t1)	
Test 7: Manual excessive temperature cut off 2 (t2)	
Test 8: Manual low temperature alarm (t3)	

Overall assessment	
	Please mark where applicable
No safety or functional deficiencies were detected	☐
No direct risk, deficiencies detected may be corrected on short term	☐
Equipment shall be taken out of operation until deficiencies are corrected	☐
Equipment does not comply – Modification / Exchange of components / Taking out of service - is recommended	☐

Comments

Date	**Signature**

13 Technical data

ASTOTHERM PLUS **REF AP220..**	**..AU** **..EU** **..UK**		**..NA**
Electrical connection	230 - 240 VAC 50 – 60 Hz		100 - 115 VAC 50 – 60 Hz
Fuses primary (F1 F2)	T4AH 250 V (5 x 20 mm)		
Fuses secondary (F3 F4)	T0630AL 250 V (TR5 type)		
Power consumption	max. 450 W		
Classification (IEC/EN 60601-1)	Class I, Type B applied part is defibrillation proof		
Classification (IEC/EN 60529)	IPX4		
Classification (MDD 93/42/EEC)	Class IIb		
Code UMDNS	10-447		
Code GMDN	47616		
Regulatory class as per FDA	II		
Dimensions Height Width Depth (incl. attachment device)	max. 145 mm 135 mm 295 mm		
Weight	2.9 kg		
Operating mode	continuous operation		
Permissible environmental conditions	Humidity	Temperature	Pressure
in operation	10% to 90% not condensing	+16°C to +32°C	700 hPa to 1060 hPa
In storage	10% to 90% not condensing	-20°C to +60°C	500 hPa to 1060 hPa
Selectable setpoint temperatures Standard setting by manufacturer	41°C 39°C 37°C		
Essential performance acc. to IEC/EN 60601-1	Regulation of the temperature of the heat exchanger to a selectable Set-temperature between 37 °C to 41°C +/- 0.5 °C		
1.Excessive temperature cut off 2. Excessive temperature cut off	42.5°C (± 0.5°C) 43.5°C (± 0.5°C)		
Low temperature alarm	$T_{Setpoint}$ – 4°C (± 0.5°C)		
Heating up time (22°C to 40°C)	approx. 1 minute		
Self-start after power interruption up to	5 seconds		

14 Compliance with international standards

Standard	Title
IEC/EN 60601-1 ANSI/AAMI ES 60601-1 CAN/CSA C22.2 No. 60601-1	Medical electrical equipment - Part 1: General requirements for basic safety and essential performance
IEC/EN 60601-1-2	Medical electrical equipment - Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility – Requirements and tests
IEC/EN 60601-1-8	Medical electrical equipment - Part 1-8: General requirements for basic safety and essential performance - Collateral Standard: General requirements, tests and guidance for alarm systems in medical electrical equipment and medical electrical systems.
ASTM F 2172-02	Standard Specification for Blood/Intravenous Fluid/Irrigation Fluid Warmers

15 Ordering information, accessories and consumables

You can order an **ASTOTHERM PLUS** Warmer using the following order numbers:

REF (Order-No.)	**Description**
AP220AU	For 4 mm infusion line, 230 - 240 VAC, Australian Plug
AP220EU	For 4 mm infusion line, 230 - 240 VAC, CEE 7/7 (Schuko) Plug
AP220NA	For 4 mm infusion line, 100 - 115 VAC, Hospital Grade Plug
AP220UK	For 4 mm infusion line, 230 - 240 VAC, Britisch Plug

Suitable Consumables:

REF (Order-No.)	**Description**
IFT40410	Sterile infusion extension made from PVC Outer diameter Ø 4 mm, length 592 cm (suitable for AP220). Filling volume 42 ml

We reserve the right to modify design and technical data without notice.

16 Guidelines and manufacturer's declaration

Guidance and manufacturer's declaration - electromagnetic emissions

ASTOTHERM PLUS is intended for use in the electromagnetic environment specified below. The customer or user of the ASTOTHERM PLUS should assure that it is used in such an environment.

Emission test	Compliance	Electromagnetic environment - guidance
RF emissions CISPR 11/EN 55011	Group 1	ASTOTHERM PLUS uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause interference in nearby electronic equipment.
RF emissions CISPR 11/EN 55011	Class A	The emissions characteristics of this equipment make it suitable for use in industrial areas and hospitals (CISPR 11 class A). If it is used in a residential environment (for which CISPR 11 class B is normally required) this equipment might not offer adequate protection to radio-frequency communication services. The user might need to take mitigation measures, such as relocating or re-orienting the equipment.
Harmonic emissions IEC/EN 61000-3-2	Class A	
Voltage fluctuations / flicker emissions IEC/EN 61000-3-3	Complies	

Guidance and manufacturer's declaration - electromagnetic immunity

The ASTOTHERM PLUS is intended for use in the electromagnetic environment specified below.
The customer or user of the ASTOTHERM PLUS should assure that it is used in such an environment.

Immunity test	Test level	Compliance level	Electromagnetic environment - guidance
Electrostatic discharge (ESD) IEC/EN 61000-4-2	± 8 kV contact ± 2 kV, ± 4 kV, ± 8 kV, ± 15 kV air	in compliance	Floors should be wood, concrete or ceramic tile. If floors are covered with synthetic material, the relative humidity should be at least 30%.
Electrical fast transient/bursts IEC/EN 61000-4-4	± 2 kV 100 kHz repetition frequency	in compliance	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC/EN 61000-4-5	± 0.5 kV, ± 1 kV Line-to-line ± 0.5 kV, ± 1 kV, ± 2 kV Line- to-ground	in compliance	Mains power quality should be that of a typical commercial or hospital environment.
Voltage dips IEC/EN 61000-4-11	0 % U_T; 0.5 cycle At 0°, 45°, 90°, 135°, 180°, 225°, 270° and 315° 0 % U_T; 1 cycle and 70 % U_T; 25/30 cycles Single phase: at 0°	in compliance	Mains power quality should be that of a typical commercial or hospital environment. If the user of the ASTOTHERM PLUS requires continued operation during power mains interruptions, it is recommended that the ASTOTHERM PLUS be powered from an uninterruptible power supply or a battery.
Voltage interruptions IEC/EN 61000-4-11	0 % U_T; 250/300 cycle	in compliance	
Rated power frequency magnetic fields IEC/EN 61000-4-8	30 A/m 50 Hz or 60 Hz	in compliance	Power frequency magnetic fields should be at levels characteristic of a typical commercial or hospital environment.

NOTE U_T is the AC mains voltage prior to application of the test level.

Guidance and manufacturer's declaration - electromagnetic immunity

The ASTOTHERM PLUS is intended for use in the electromagnetic environment specified below.
The customer or user of the ASTOTHERM PLUS should assure that it is used in such an environment.

Immunity test	Test level	Compliance level	Electromagnetic environment – guidance Recommended separation distance
Conducted disturbances induced by RF fields IEC/EN 61000-4-6	3 Vrms 0.15 MHz to 80 MHz 6 Vrms in ISM bands between 0.15 MHz and 80 MHz 80 % AM at 1 kHz	in compliance	$d = 1.2\sqrt{P}$
Radiated RF EM fields IEC/EN 61000-4-3	3 V/m 80 MHz to 2.7 GHz 80 % AM at 1 kHz	in compliance	$d = 1.2\sqrt{P}$ 80 MHz to 800 MHz $d = 2.3\sqrt{P}$ 800 MHz to 2.7 GHz

Portable and mobile RF communications equipment should be used no closer to any part of the ASTOTHERM PLUS, including cables, than the recommended separation distance calculated from the equation applicable to frequency of the transmitter.

Where P is the maximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer and "d" is the recommended separation distance in meters (m).
Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, [a] should be less than the compliance level in each frequency range [b].
Interference may occur in the vicinity of equipment marked with the following symbol:

NOTE1. At 80 MHz and 800 MHz, the higher frequency range applies.
NOTE2. These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

[a] Field strengths from fixed transmitters, such as base stations for radio (cellular/cordless) telephones and land mobile radios, amateur radio, AM and FM radio broadcast and TV broadcast cannot be predicted theoretically with accuracy. To assess the electromagnetic environment due to fixed RF transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the ASTOTHERM PLUS is used exceeds the applicable RF compliance level above, the ASTOTHERM PLUS should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as reorienting or relocating the ASTOTHERM PLUS.
[b] Over the frequency range 150 kHz to 80 MHz, field strength should be less than 3 V/m.

Recommended separation distances between
Portable and mobile RF communications equipment and the ASTOTHERM PLUS

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

Rated maximum output power of transmitter (W)	Separation distance according to frequency of transmitter (m)		
	150 kHz to 80 MHz $d = 1.2\sqrt{P}$	80 MHz to 800 MHz $d = 1.2\sqrt{P}$	800 MHz to 2.7 GHz $d = 2.3\sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.38	0.38	0.73
1	1.2	1.2	2.3
10	3.8	3.8	7.3
100	12	12	23

For transmitters rated at maximum output power not listed above, the recommended separation distance d in meters (m) can be estimated using the equation applicable to the frequency of the transmitter, where P is the maximum output power rating of the transmitter in Watts (W) according to the transmitter manufacturer.
NOTE 1: The compliance level between 80 MHz and 2.7 GHz is intended to decrease the likelihood that mobile/portable communication equipment could cause interference if it is inadvertently brought into patient areas. For this reason, an additional factor of 10/3 is used in calculating the recommended separation distance for transmitters in this frequency range.

NOTE 2: These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.