HUNTLEIGH IRT10

Anwendungshinweise

Kullanım Talimatları

Brugsveiledning

Instrucciones de uso

使用方

Mode d'emploi

Bruksanvisning

Gebruiksaanwijzing

aanwijzing

; χρήσης

INSTRUCTIONS FOR USE

Bruksa

alimatları

使用方法

Käyttöohjeet

Instruções de Utilização

Istruzioni per l'uso

Anwendungshinweise

Οδηγίες χρήσης

Anwendungshinweise

Wireless Tympanic Thermometer

	onte		
1.		al Safety Information	
		Varnings / Cautions	
2.	Introd	uction	6
	2.1	Features of the IRT10 Wireless Themometer	6
	2.2	Intended Use	
	2.2	Contraindications	
_	2.4	About This Manual	7
3.	Descri	iption of Controls, Indicators, Symbols and	
	Displa	ys	8
	3.5	Thermometer Controls and Display	8
	3.2	Description of Symbols / Indicators	
4.	Setup		11
	4 .1	Unpacking and Inspection	
	4.2	List of Components	
	4.3	Connecting the Thermometer to the host	11
	4.4	Installing the Batteries	
_	4.5	Pairing the Thermometer and Main Unit	
5.	Using	the Thermometer	13
	5.1	Temperature Measurement	13
6.	Care 8	& Cleaning	15
	6.1	General	
	6.2	Returning the IRT10 Thermometer and System Components	
	6.3	Service	
	6.4	Periodic Safety Checks	
7.	Troubl	leshooting	16
	7.1	General	_
_	7.2	Obtaining Technical Assistance	
B.	Electr	omagnetic Compatibility	17
9.	Specif	ications	21
	9.1	Equipment Classification	21
	9.2	Standards	
	9.3	General	
	9.4	Environmental	
	9.5	Measurement Parameters	
10	. Warra	anty & Service	23
	10 1	Sanica Paturas	22

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Revision History

The documentation part number and revision number indicate its current edition. The revision number changes when a new edition is printed in accordance with the revision history of the documentation. Minor corrections and updates which are incorporated at reprint do not cause the revision number to change. The document part number changes when extensive technical changes are incorporated.

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The IRT10 is in conformity with the Medical Devices Directive 93/42/EEC as amended by 2007/47/EC and has been subject to the conformity assurance procedures laid down by the Council Directive.

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1. General Safety Information

This section contains important safety information related to general use of the IRT10 wireless thermometer. Other important safety information appears throughout the manual



Important! Before using this equipment, please study this manual carefully and familiarise yourself with the controls, display features and operating techniques. Ensure each user fully understands the safety and operation of the unit, as misuse may cause damage to the unit or injury to the user or patient.

1.1 Warnings / Cautions



General warning / caution



Refer to Instructions for Use



WARNING: Check the equipment prior to use and ensure its safe and proper use.



WARNING: Before use, carefully read directions for use, including all warnings, cautions, and instructions.



WARNING: If the battery shows any signs of damage, leakage, or cracking, it must be replaced immediately, by a qualified service person, and only with a battery approved by the manufacturer.



WARNING: The monitor is intended only as an adjunct in patient assessment. It must be used in conjunction with clinical signs and symptoms.



WARNING: It is possible that any radio frequency transmitting equipment and other sources of electrical noise such as cellular phones, due to close proximity or strength of a source, may result in disruption of performance.



CAUTION: U.S. Federal law restricts this device to sale by or on the order of a licensed healthcare practitioner.

2. Introduction



WARNING: The Wireless Tympanic Thermometer is intended only as an adjunct in patient assessment. It must be used in conjunction with clinical signs and symptoms.

This manual contains information about the Wireless Tympanic Thermometer (Order code: IRT10).

It is available as an option for the SC300 SPOT CHECK Vital Signs Monitor.





2.1 Features of the IRT10 Wireless Themometer

Physical

The IRT10 wireless tympanic thermometer is a clinical grade hand held thermometer indicated for the intermittent measurment of body temperature in patients of all ages.

Electrical

The Thermometer is powered by two AAA dry cells.

Display

The Thermometer presents the following information on the integrated display:

- Battery status
- Lens cover indicator
- Wireless pairing status
- Temperature measurment
- Measurement scale °C or °F

2.2 Intended Use

The Thermometer is intended for use by trained healthcare professionals in healthcare settings to measure human body temperature in patients of all ages. It uses an infra red technique to measure the amount of infrared energy reflected from the tympanic membrane.

2.2 Contraindications

The Thermometer should not be used if the following situations are present:

Do not use the thermometer if there is any blood or drainage in the ear canal.

Do not use the thermometer if the patient presents with inflamatory condition of the ear canal.

Do not use the thermometer if the ear canal is blocked with cerumen.

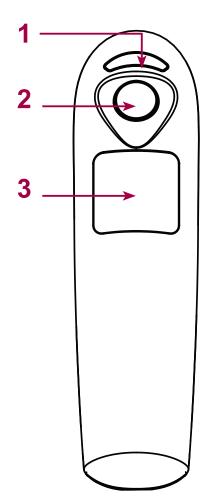
Do not use the thermometer if ear drops or medication has been applied to the ear

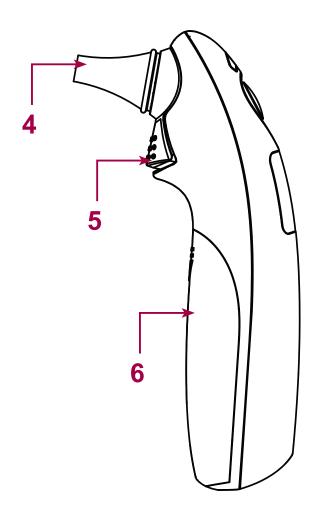
2.4 About This Manual

Read the entire manual including the Safety Information section, before you operate the monitor.

3. Description of Controls, Indicators, Symbols and Displays

3.5 Thermometer Controls and Display





1	Temperature probe status indicator	
2	Start measurment	
3	Display	
4	Infra red sensor	
5	Lens ejector	
6	Battery compartment	



Key	Function / Display	
	Battery status	
嶽	Lens filter status	
3	Wireless status	
°C°F	Scale °C or °F	
188.8	Measurement	

3.2 Description of Symbols/ Indicators

The symbols and Indicators are described as follows:

Symbols	Description	
<u> </u>	Attention, consult accompanying documents.	
	Battery charging indicator lamp	
(3)	Refer to instructions for use	
IPX1 Protection against vertically falling water drops		
Date of manufacture		
SN Serial number		

Symbols	Description		
(E 0088	Conformité Européenne Complies with medical device directive 93/42/ EEC		
Disposal instructions			
	Manufacturer		
<u> </u>	This way Up		
Fragile – Handle with care			
Limit of stacking layers			
	Keep dry		

4. Setup



WARNING: The IRT10 Wireless Thermometer is a prescription device and is to be operated by qualified personnel only. It is designed for use by medical clinicians.



WARNING: Discarded battery may explode during incineration. Follow local government ordinances and recycle instructions regarding disposal or recycling of device components, including batteries. Do not dispose of batteries in refuse containers.

4.1 Unpacking and Inspection

The Thermometer is shipped in one carton. Examine the carton carefully for evidence of damage. Contact Huntleigh Healthcare Ltd Service Department immediately if any damage is discovered. Return all packing material and Thermometer. Refer to the Maintenance section for instructions on returning damaged items.

4.2 List of Components

Quantity	Item	Quantity	Item
1	Thermometer	1	Lens covers
2	AAA dry cells	1	Cradle

4.3 Connecting the Thermometer to the host

To facilite ease of use, the Thermometer can be fitted to the SC300 using the cradle supplied.





4.4 Installing the Batteries



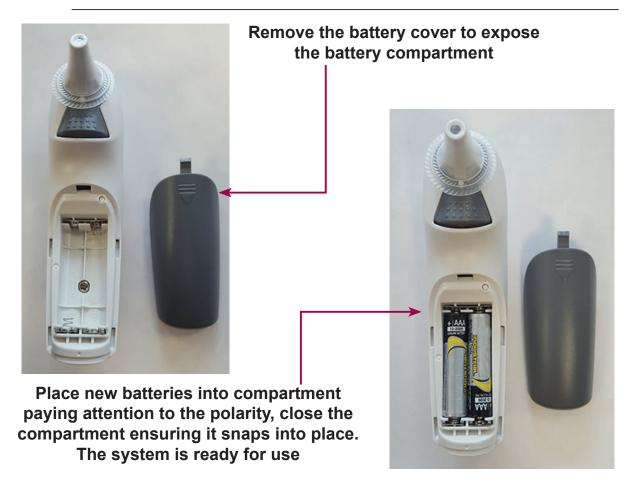
WARNING: Dispose of Battery in accordance with local requirements and regulation. Follow local instructions regarding disposal or recycling of batteries.



CAUTION: Measured or displayed data may not be assured in the low battery or the critical low battery condition.



CAUTION: Discarded battery may explode during incineration. Recycle used batteries properly. Do not dispose of batteries in refuse containers.



4.5 Pairing the Thermometer and Main Unit

- 1. Make sure the main unit and ear thermometer are both switched OFF.
- 2. Press and hold the "ejector key" on the thermometer and switch it ON.
- 3. The thermometer display will alternate between °C and °F and then display "SE".
- 4. When "SE" is displayed, release the ejector button and switch the main unit ON.
- 5. Pairing will be established when the icon is displayed on the host unit.
- 6. If pariring has failed, no icon will be displayed.

5. Using the Thermometer



The Thermometer must be calibrated at least once every two years, contact your service personnel when calibration is necessary



Use only specified protective lens covers



The protective lens cover is single-use. Repeated use might give rise to cross infection.



The protective lens cover must be used when measuring, if not, it might cause cross infection or inaccurate readings.



Before use, check whether the cover is free from damage, if not, don't use.



Handle the thermometer with care, it should be stored in the cradle when not in use.



Discard the protective lens cover in accordance with the local regulations.



During the monitoring process, the temperature measuring instrument will automatically check itself once per hour. Self-checking will last 2 seconds, and will not affect the normal working of the temperature monitor.

Measurments are sent via a wireless connection established between the thermometer and the main unit.

Both items (Thermometer and unit) must be paired to enable data transfer.

5.1 Temperature Measurement

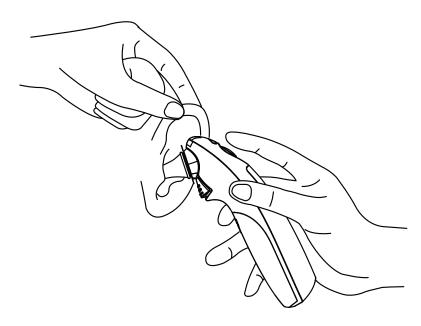
- 1. Install a new protective lens cover to the thermometer.
- 2. When correctly fitted, the



icon changes to



3. Position the patient with their head moved to one side, gently pull the ear backward to straighten the ear canal. Gently place the sensor into the ear canal.



- 4. Press the START button once, the sensor will sound a quick double beep indicating that its ready to measure.
 - Press the START button once again and after a short period, the sensor will sound a long single tone indicating that the measurement is complete.
- 5. Remove the thermometer.
- 6. The measurement will be shown on the thermometer's screen, it will be automatically transferred to the monitor.



Thermometer display



Monitor display

- 7. Press the eject key to remove the disposable lens cover.
- 8. Place the lens cover in the appropriate waste collection point and place the thermometer onto the unit

6. Care & Cleaning

6.1 General

The probe tip is the most delicate part of the thermometer. It has to be clean and intact to ensure accurate readings.

Gently wipe the surface of the probe tip with a cotton swab or soft cloth moistened with alcohol.

After the alcohol has completely dried, fit a new lens filter and take a temperature measurement.

If the probe tip is damaged, please contact your service agent.

Use a soft, dry cloth to clean the thermometer display and exterior.

Do not use abrasive cleaners.

Never submerge the thermometer in water or any other liquid.

Store thermometer and lens filters in a dry location free from dust and contamination and away from direct sunlight.

6.2 Returning the IRT10 Thermometer and System Components

Contact Huntleigh Healthcare Ltd Service Department for shipping instructions. Pack the accessory items in its original shipping carton. If the original carton is not available, use a suitable carton with appropriate packing material to protect the device during shipping. Return by any shipping method that provides proof of delivery.

6.3 Service

Apart from a 2 year calibration check, the Wireless Thermometer requires no routine service other than cleaning or battery maintenance that is mandated by the user's institution. For more information, refer to Smartsigns® Compact 300 service manual. Qualified service personnel in the user's institution should perform periodic inspections of the equipment. If service is necessary, contact qualified service personnel or Huntleigh Healthcare Ltd Service Department.

If the institution's service personnel cannot correct problems, the equipment should be returned to Huntleigh Healthcare Ltd for service. Contact Huntleigh Healthcare Ltd Service Department for return instructions.

6.4 Periodic Safety Checks

It is recommended that the following checks be performed every 24 months.

- Inspect the equipment for mechanical and functional damage.
- Inspect the safety relevant labels for legibility.

7. Troubleshooting



WARNING: If you are uncertain about the accuracy of any measurement, check the patient's vital signs by alternate means; then make sure the monitor is functioning correctly.



WARNING: Only qualified service personnel should remove the cover. There are no user-serviceable parts inside.

7.1 General

If the equipment is unable to perform any of its monitoring functions because of the loss of software control or a detected hardware malfunction, an error code is presented. In the unlikely event of the unit developing a fault, fault codes will be shown in the corresponding area and the related parameters will flash on the screen

Description	Cause	Solution	
Low power Low power		Replace batteries	
	Auto shutdown	Restart the device	
Blank Screen	Battery installation incorrectly	Check the correct installation of the batteries	
	Low power	Replace batteries	
	The screen is still blank	Contact service personnel	
Er1	The environment temperature is out of range 10~40°C (33.8-39.2°F)	Move the device to appropriate environment and wait 30 minutes prior to taking a temperature.	
Er2 The environment temperature is not stable		Do not take measurement until the environment temperature is stable	
Er3	Infrared module failure	Contact service personnel	
Hi The measurement is out of range > 42.2°C (108°F)		Make sure the operation method is correct and the lens filter is new and clean. Remeasure.	
Lo The meaurment is out of range <34°C (108°F)		Make sure the operation method is correct and the lens filter is new and clean. Remeasure.	

7.2 Obtaining Technical Assistance

For technical information and assistance, or to order a service manual, call Huntleigh Healthcare Ltd Service Department. The service manual includes information required by qualified service personnel when servicing the Smartsigns® Compact 300.

When calling the Huntleigh Healthcare Ltd Service Department, you may be asked to tell the representative the software version number of your equipment. Qualified service personnel or Huntleigh Healthcare Ltd Service Department may help you check the software version installed in your equipment.

8. Electromagnetic Compatibility

Make sure the environment in which the equipment is installed is not subject to strong sources of electromagnetic interference (e.g. radio transmitters, mobile phones).

This equipment generates and uses radio frequency energy. If not installed and used properly, in strict accordance with the manufacturer's instructions, it may cause or be subject to interference. Type-tested in a fully configured system, complies with EN60601-1-2, the standard intended to provide reasonable protection against such interference. Whether the equipment causes interference may be determined by turning the equipment off and on. If it does cause or is affected by interference, one or more of the following measures may correct the interference:

- Reorienting the equipment
- Relocating the equipment with respect to the source of interference
- Moving the equipment away from the device with which it is interfering
- Plugging the equipment into a different outlet so that the devices are on different branch circuits



WARNING: The use of accessories, transducers and cables other than those specified, with the exception of transducers and cables sold by the manufacturer of the Smartsigns® Compact 300 as replacement parts for internal components, may result in increased emissions or decreased immunity of the Smartsigns® Compact 300.



WARNING: The Smartsigns® Compact 300 should not be used adjacent to or stacked with other equipment and that if adjacent or stacked use is necessary, the Smartsigns® Compact 300 should be observed to verify normal operation in the configuration in which it will be used



WARNING: 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 Smartsigns® Compact 300 including cables specified by the manufacturer. Otherwise, degradation of the performance of this equipment could result.

Guidance and Manufacturer's declaration - electromagnetic emissions

The IRT10 infrared ear thermometer is intended for use in the electromagnetic environment specified below. The customer or the user of the IRT10 infrared ear thermometer should assure that it is used in such an environment.

Emissions Test	Compliance	Electromagnetic Environment - guidance
RF emissions CISPR 11	Group 1	The IRT10 infrared ear thermometer uses RF energy only for its internal function. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment.
RF emissions CISPR 11	Class A	
Harmonic emissions IEC 61000-3-2	Class A	The IRT10 infrared ear thermometer is suitable for use in all establishments, other than domestic and those directly connected to the public low-voltage power supply network
Voltage fluctuations / flicker emissions IEC 61000-3-3	Complies	that supplies buildings used for domestic purposes.

Guidance and Manufacturer's declaration - electromagnetic immunity

The IRT10 infrared ear thermometer is intended for use in the electromagnetic environment specified below. The customer or the user of the IRT10 infrared ear thermometer should assure that it is used in such an environment.

Immunity Test	IEC 60601 test level	Compliance level	Electromagnetic Environment - guidance
			Portable and mobile RF communications equipment should be used no closer to any part of the Smartsigns® Compact 300, including cables, than the recommended separation distance calculated from the equation applicable to the frequency of the transmitter.
Conducted RF IEC 61000-4-6	3 Vrms 150kHz to 80MHz outside ISM bands ^a 6 Vrms 150 kHz to 80 MHz in ISM and amateur radio bands	3V	$d = 1.2 \sqrt{P}$
Radiated RF IEC 61000-4-3	3 V/m 80MHz to 2.5GHz	3V/m	$d = 1.2 \sqrt{P}$ $d = 2.3 \sqrt{P}$ 80MHz to 800MHz $d = 2.3 \sqrt{P}$ 800MHz to 2.5GHz
			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 metres (m). ^b Field strengths from fixed RF transmitters, as determined by an electromagnetic site survey, ^c should be less than the compliance level in each frequency range ^d . Interference may occur in the vicinity of the equipment marked with the following symbol: $\left(\left(\begin{pmatrix} \bullet \end{pmatrix}\right)\right)$

NOTE 1 At 80MHz and 800MHz, the higher frequency range applies.

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

- ^a The ISM (industrial, scientific and medical) bands between 150 kHz and 80 MHz are 6,765 MHz, to 6,795 MHz; 13,553 MHz to 13,567 MHz; 26,957 MHz to 27,283 MHz; and 40,66 MHz to 40,70 MHz.
- The compliance levels in the ISM frequency bands between 150 kHz and 80 MHz and in the frequency range 80 MHz to 2,5 GHz are intended to decrease the likelihood that mobile/portable communications 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 these frequency ranges.
- 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 IRT10 infrared ear thermometer is used exceeds the applicable RF compliance level above, the IRT10 infrared ear thermometer should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the Smartsigns® Compact 300.
- Over the frequency range 150kHz to 80kHz, field strengths should be less than 3V/m.

Guidance and Manufacturer's declaration - electromagnetic immunity

The IRT10 infrared ear thermometer intended for use in the electromagnetic environment specified below. The customer or the user of the IRT10 infrared ear thermometer should assure that it is used in such an environment.

Immunity Test	IEC 60601 test level	Compliance level	Electromagnetic Environment - guidance
Electrostatic discharge (ESD)	± 6 kV contact	± 6 kV contact	Floors should be wood, concrete or ceramic tile. If floors are covered with
IEC 61000-4-2	± 8 kV air	± 8 kV air	synthetic material, the relative humidity should be at least 30%.
Electrical fast transient burst	± 2 kV for power supply lines	± 2 kV for power supply lines	Mains power quality should be that of a typical commercial or hospital environment.
IEC 61000-4-4	± 1 kV for input/ output lines	± 1 kV for input/ output lines	
Surge	± 1 kV line(s) to line(s)	± 1 kV line(s) to line(s)	Mains power quality should be that of a typical commercial or hospital environment.
IEC 61000-4-5	± 2 kV line(s) to earth	± 2 kV line(s) to earth	
Voltage dips, short interruptions and voltage variations on power supply input	$<5 \% U_{\rm r}$ (>95 % dip in $U_{\rm r}$) for 0,5 cycles	$<5 \% U_{\rm r}$ (>95 % dip in $U_{\rm r}$) for 0,5 cycles	Mains power quality should be that of a typical commercial of hospital environment. If the user of the IRT10 infrared ear thermometer requires continued operation
lines IEC 61000-4-11	$40 \% U_{\rm r} \ (60 \% {\rm dip \ in} \ U_{\rm r}) \ {\rm for \ 5 \ cycles}$	$40 \% U_{\rm r}$ (60 % dip in $U_{\rm r}$) for 5 cycles	during power mains interruptions, it is recommended that the IRT10 infrared ear thermometer is powered from an uninterruptible power supply or battery,
	70 % $U_{\rm r}$ (30 % dip in $U_{\rm r}$) for 25 cycles	70 % $U_{\rm r}$ (30 % dip in $U_{\rm r}$) for 25 cycles	by specifying the battery option at time of purchase.
	<5 % $U_{\rm r}$ (>95 % dip in $U_{\rm r}$) for 5 s	<5 % $U_{\rm r}$ (>95 % dip in $U_{\rm r}$) for 5 s	
Power frequency (50/60Hz) magnetic field	3 A/m	3 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial or hospital environment.
IEC 61000-4-8			

NOTE U_r is the a.c. mains voltage prior to the application of the test level.

Recommended separation distances between portable and mobile RF communications equipment and the Smartsigns® Compact 300

The IRT10 infrared ear thermometer is intended for use in an electromagnetic environment in which radiated RF disturbances are controlled. the customer or user of the IRT10 infrared ear thermometer can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the IRT10 infrared ear thermometer as recommended below, according to the maximum output power of the communications equipment.

Rated maximum output power of	Separation distance according to frequency of transmitter m			
transmitter	150kHz to 80MHz	80MHz to 800MHz	800MHz to 2.5GHz	
W	$d = 1.2\sqrt{P}$	$d = 1.2\sqrt{P}$	$d = 2.3 \sqrt{P}$	
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 a maximum output power not listed above, the recommended separation distance d in metres (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 At 80MHz and 800MHz, the separation distance for the higher frequency range applies. NOTE 2 These guidelines may not apply in all situations. Electromagnetic propagation is affected by absorption and reflection from structures, objects and people.

9. Specifications

9.1 Equipment Classification

Type of protection against electric shock.	Internally powered equipment
Degree of protection against electric shock	BF
Mode of operation.	Continuous
Degree of protection against harmful ingress of particles and/or water.	IPX1
Degree of safety of application in the presence of a flammable anaesthetic	Equipment not suitable for use in the presence of a FLAMMABLE ANAESTHETIC MIXTURE WITH AIR, OXYGEN OR NITROUS OXIDE

9.2 Standards

MDD 93/42/EEC	EN ISO13485:2012+AC2012
EN ISO14971: 2012	EN 60601-1: 2006/ AC:2013
EN 60601-1-2: 2007/AC:2010	EN60601-1-6:2010
EN 980:2008	EN 1041: 2008
EN ISO10993-1:2009	EN ISO10993-5:2009
EN ISO 10993-10:2010	EN 1060-1:1995+A2:2009
EN 1060-3:1997+A2:2009	EN 1060-4:2004
EN ISO 80601-2-30:2009+A1:2013	EN ISO 80601-2-61:2011
EN 62366:2008	EN62304:2006

9.3 General

Battery	2 x Type LR03 AAA1.5VDC
Battery Life	3000 measurements
Size	133 x 63.5 x 36.4 mm
Weight	80g
Automatic Shut down	60s±10s
Service Life	7 years

9.4 Environmental

Operation		
Temperature	15 °C to 36 °C (59 °F to 96.8 °F)	
Humidity	≤85% non-condensing	
Altitude	700 hPa to 1060hPa	
Transport and Storage		
Temperature	–25 °C to 55 °C (-13 °F to 131 °F)	
Humidity	≤85% non-condensing	
Altitude	700 hPa to 1060 hPa	

Note: The system may not meet its performance specifications if stored or used outside the manufacturer's specified temperature and humidity range.

9.5 Measurement Parameters

Probe Type	Tympanic Infrared
Range	34 °C to 42.2 °C (93.2 °F to 108°F)
Accuracy	35° C to 42° C (95 °F to 107.6°F): ±0.2° C (±0.4° F) Other Range: ±0.3° C (±0.5° F)
Resolution	0.1°C (0.1 °F)
Measurement Interval	≥12s
Masurement duration	3s±1s

10. Warranty & Service

Huntleigh Healthcare Diagnostic Products Division standard terms and conditions apply to all sales. A copy is available on request. These contain full details of warranty terms and do not limit the statutory rights of the consumer.

10.1 Service Returns

If for any reason the Wireless Thermometer has to be returned, please:

- Clean the product following the instructions in this manual.
- Pack it in suitable packing.
- Attach a decontamination certificate (or other statement declaring that the product has been cleaned) to the outside of the package.
- Mark the package 'Service Department'

For further details, refer to NHS document HSG(93)26 (UK only).

Huntleigh Diagnostics reserve the right to return product that does not contain a decontamination certificate.

A service manual is available for the Smartsigns® series. It contains service information, parts lists and fault finding guidelines. The service manual can be obtained by contacting your local supplier or:-

Service Department. Huntleigh Healthcare, Diagnostic Products Division, 35, Portmanmoor Rd., Cardiff. CF24 5HN United Kingdom.

Tel: +44 (0)29 20485885 Fax: +44 (0)29 20492520

Email: sales@huntleigh-diagnostics.co.uk

service@huntleigh-diagnostics.co.uk www.huntleigh-diagnostics.com



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