

MASiMO Stork Replacement Sensor User Manual

Home » MASiMO stork Replacement Sensor User Manual



Contents

- 1 MASiMO Stork Replacement
- Sensor
- **2 Product Information**
- **3 Product Usage Instructions**
- **4 DESCRIPTION**
- **5 INSTRUCTIONS**
- **6 CLEANING**
- **7 SAFETY PRECAUTIONS**
- 8 Documents / Resources



MASiMO Stork Replacement Sensor



Product Information

Product Name: Replacement SensorManufacturer: Masimo Corporation

· Application Site: Foot

· Age/Baby Size: 0-18 months

• Measurements: 2.5 (L) x 1.9 (W) x .03 (H) inches (6.35 cm x 4.83 cm x 0.76 cm)

• Weight: 0.03 lbs. (13g)

• Battery Type: Li-ion rechargeable

• Battery Run Time: 16 hours*

· Charging Time: 2 hours

• SpO2 Accuracy, No Motion, ARMS 1: 1.5%

• SpO2 Accuracy, Motion, ARMS 2: (not provided)

• Storage Temperature: (not provided)

• Operating Temperature: (not provided)

• Storage/Transport Humidity: (not provided)

• Operating Humidity: (not provided)

• Atmospheric Pressure: (not provided)

Product Usage Instructions

- 1. Charge the sensor:
 - 1. Plug in the USB-C cable to the 5W power adapter.
 - 2. Plug in the power adapter into a wall socket.
 - 3. Peel and stick the cord clip to secure the sensor charge cord.
 - 4. Connect the magnetic charger to the sensor.
 - 5. The sensor is fully charged when the LED light turns solid white.

- 2. Insert the Sensor into the Boot:
 - 1. Insert the sensor into the boot at an angle, toe end first, with the Masimo logo facing down and press down to secure the sensor. The sensor should sit flush in the boot.
 - 2. Follow the instructions from your Stork system, boot, and/or in the app to ensure the boot fits properly and the straps are not too tight to avoid injury.
- 3. Remove the Sensor:
 - 1. Gently push the sensor through the opening on the bottom of the boot to remove the sensor from the boot.
- 4. Cleaning Caution:
 - 1. Remove the sensor from the baby and the boot.
 - 2. Clean the sensor by wiping it with a 70% isopropyl alcohol pad or mild detergent.
 - 3. Allow the sensor to dry completely prior to charging or use.
- 5. Pair the Sensor with your device (phone not included):
 - 1. Open the Stork App and click on Sensor.
 - 2. Follow the on-screen instructions to pair the sensor.

If you need further assistance or have any questions, please visit the Stork support page: www.masimostork.com/en-us/support/contact-us.html

Replacement Sensor

Not intended for use as a medical device or to replace a medical device. Do not use it to diagnose, cure, treat, alleviate, or prevent any disease or health condition.

Download the Stork App before using this product. For help, go to www.masimostork.com for additional tips and tutorials, a full list of supported devices, warranty, trouble shooting, and customer support. Follow the in app instructions for device pairing and general setup.

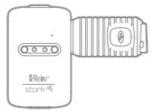
DESCRIPTION

StorkTM is a smart home baby monitoring system designed for parents to monitor a healthy baby at home. The Stork Sensor is a wireless sensor that fits into the bottom of the Stork Boot. It uses Masimo SET® technology to track baby's health data, including oxygen saturation, pulse rate, and temperature. It is also capable of determining baby's position and notifying you in-app if baby is face up or face down.

Use as a Stork replacement sensor or as an extra sensor that will allow you to continuously track your baby's health data without charging downtime.

Contents:

Stork Sensor



Magnetic Charge Cable & Cord Clip



• 5W Power Adapter



Not included: Boot, Boot Straps, Camera, or Hub (all sold separately).

INSTRUCTIONS

Charge the sensor

CAUTION: Make sure the sensor is fully changed before use.

- 1. Refer to Fig. 1. Plug in the USB-C cable to the 5W power adapter.
- 2. Refer to Fig. 2. Plug in the power adapter into a wall socket.
- 3. Refer to Fig. 3. Peel and stick the cord clip to secure the sensor charge cord.
- 4. Refer to Fig. 4. Connect the magnetic charger to the sensor.
- 5. Refer to Fig. 5. The sensor is fully changed when the LED light turns solid white.

Pair the Sensor with your device

(phone not included)

- 1. Refer to Fig. 6. Open the Stork App and click on Sensor.
- 2. Follow the on screen instructions to pair the sensor.

Insert the Sensor into the Boot

- 1. Refer to Fig. 7. Insert the sensor into the boot at an angle, toe end first, with the Masimo logo facing down and press down to secure the sensor. The sensor should sit flush in the boot.
- 2. Refer to Fig. 8.

WARNING: Follow the instructions from your Stork system, boot and/or in the app to ensure the boot fits properly and the straps are not too tight to avoid injury

Remove the Sensor

1. Refer to Fig. 9. Gently push the sensor through the opening on bottom of the boot to remove the sensor from the boot.

CLEANING

CAUTION:

- Do not use undiluted bleach or any cleaning solution other than those recommended here because permanent damage to the sensor could occur.
- Do not immerse the connector of the sensor or cable in any liquid solution.
- Using excessive force when removing the saensor may damage it.

To clean the sensor

- 1. Remove the sensor from the baby and the boot.
- 2. Refer to Fig. 10. Clean the sensor by wiping it with a 70% isopropyl alcohol pad or mild detergent.
- 3. Allow the sensor to dry completely prior to charging or use

CUSTOMER SUPPORT

For product support, along with troubleshooting for your Stork product, please go to the Stork support page: www.masimostork.com/en-us/support/contact-us.html

SAFETY PRECAUTIONS

- Stork is not intended for use as a medical device or to replace a medical Do not use it to diagnose, cure, treat, alleviate, or prevent any disease or health condition.
- Do not self-diagnose or self-medicate on the basis of the Always consult your doctor.
- Periodically check the areas where the user's skin contacts with the sensor to prevent potential issues, since there are skin-contacting components, there is a risk of skin irritation, pressure injury, and general discomfort while a sensor is applied to a
- Before using Stork, read the safety precautions carefully. Always consult a physician if you have concerns
 about your baby's well-being. Call emergency services if you believe your baby is having a medical
- Do not rely on Stork for a clinical Stork is not intended to be used as a medical device or to replace a physician or health care professional. A clinical assessment of your baby should be done by a physician. For safe use, do not use any component of the Stork System if it appears damaged.
- Do not adjust, repair, open, disassemble, or modify Stork. Such changes may lead to injury and/or incorrect
- · Misapplied sensors may cause incorrect
- Do not leave the sensor components unattended, small items may become choking
- · Carefully position any cables to avoid
- Remove the Stork Sensor before bathing to prevent
- Avoid strapping the boot too tightly around the foot to avoid
- To avoid skin injury, consider alternating feet after it has been used for more than 12 hrs.
- Only use the AC power supply and cable included with your Stork Sensor to prevent damage to the
- Do not monitor more than one person at a time with Stork.
- The Stork Sensor is to only be used with Masimo Stork authorized
- Do not use the Stork Sensor if it has visible defects, appears damaged, or seams Otherwise the sensor may not work properly.
- Check the sensor for proper securement and alignment frequently and adjust the sensor as necessary.
- Do not use additional tape to secure the sensor to the site. This can restrict blood flow and cause incorrect readings. Use of additional tape can also cause skin damage, and/or pressure injury or damage the sensor.

- Avoid placing the sensor directly under bright lights (e.g. fluorescent lights, infrared heating lamps, and direct sunlight) as they can interfere with the performance of the Cover the sensor site with opaque material, if required.
- · Check pairing before use to ensure proper wireless
- To prevent damage, do not soak or immerse the sensor in any liquid
- Do not modify or alter the sensor in any way. Alteration or modification may affect

Things that interfere with the light can affect your SpO2 accuracy. Some things can be controlled, while others need awareness:

Controllable things:

- · Keeping feet dry and free of foreign objects.
- · Not blocking the light through your foot.
- Avoiding direct exposure to bright lights, including direct sunlight.
- · Keeping the foot still.
- Warm the baby's foot to improve circulation.
- Avoid use on the same leg with blood pressure cuff inflated.
- Keep away from other electrical equipment that may cause readings to be affected (e.g., microwaves, strong radio transmitters).

Other things that require awareness:

- Skin pigment or color.
- · Skin or foot thickness.
- Skin conditions that may affect the ability to comfortably wear theboot (e.g.: eczema).
- Foot deformities (e.g. extreme club foot).
- Health conditions affecting how oxygen is carried in your blood (e.g., sickle cell, severe anemia).
- · Poor blood circulation.
- Presence of blood components not able to carry oxygen (e.g., elevated carbon monoxide levels in the blood.
- · Age and medical history.

SENSOR STATUS LIGHT

Light Indicator Color	What does it mean?
Blinking Green	Sensor is on and waiting to be paired.
Solid Blue	Sensor is paired.
Blicking Orange	Sensor battery is low.
Blinking Red	Sensor battery is very low/depleted.
Blinking White	Sensor battery is charging.
Solid White	Sensor battery is fully charged.
Blinking Red in a pattern to indicate a number code	Sensor has a problem (non-battery related)

BATTERY

Battery type:	Li-ion rechargable	Battery run time:	16 hours*	Charging Time:	2 hours
---------------	--------------------	-------------------	-----------	----------------	---------

Minimum of 16 hours in typical continuous usage until fully discharged

ENVIRONMENTAL

Storage Temperature	-20°C to + 60°C @ ambient humidity
Operating Temperature	5°C to + 35°C @ ambient humidity
Storage/Transport Humidity	10-95% RH (non-condensing) @ ambient temperature
Operating Humidity	10-95% RH (non-condensing) @ ambient temperature
Atmospheric Pressure	540 to 1060 mBar at ambient temperature and humidity

PERFORMANCE SPECIFICATIONS

The Stork Sensor has the following specifications:

Stork Sensor			
Application Site:	Foot		
Age/Baby Size:	0-18 months		
Measurements:	2.5" (L) x 1.9" (W) x .03" (H) (6.35 cm x 4.83 cm x 0.76 cm		
Weight:	0.03 lbs. (13g)		
Oxygen level SpO2 performance (Arms) specification	ons (70-100%)SaO2		
SpO2 Accuracy, No Motion, Arms 1	1.5 %		
SpO2 Accuracy, Motion, Arms 2	1.5 %		
SpO2 Accuracy, Low Perfusion, Arms 3	2 %		
Pulse Rate (Arms) specifications from 25-240 BPM			
Pulse Rate No Motion, Arms 4	≤ 3 BPM		
Pulse Rate, Motion, Arms	≤ 5 BPM		
Pulse Rate, Low Perfusion, Arms	≤ 3 BPM		
Temperature Measurement Accuracy			
±0.3°C (±0.54°F) in the range of 25°C to 43°C (77°F to 109.4°F)			
Position			
The sensor will distinguish if the baby changes postion to face down.			

Note: Arms accuracy is a statistical calculation of the difference between device measurements and reference measurements. Approximately two-thirds of the device measurements fell within +/- Arms of the reference measurements in a controlled study.

- 1. The Masimo SET Technology has been validated for no motion accuracy in human blood studies on healthy adult male and female volunteers with light to dark pigmented skin in induced hypoxia studies in the range of 70–100% SpO2 against a laboratory co-oximeter.
- 2. The Masimo SET Technology has been validated for motion accuracy in human blood studies on healthy adult male and female volunteers with light to dark pigmented skin in induced hypoxia studies while performing rubbing and tapping motions, at 2 to 4 Hz at an amplitude of 1 to 2 cm and a non-repetitive motion between 1 to 5 Hz at an amplitude of 2 to 3 cm in induced hypoxia studies in the range of 70–100% SpO2 against a laboratory co-oximeter.

- 3. The Masimo SET Technology has been validated for low perfusion accuracy in bench top testing against a Biotek Index 2 simulator and Masimo's simulator with signal strengths of greater than 0.02%.
- 4. The Masimo SET Technology has been validated for pulse rate accuracy for the range of 25-240 bpm in bench top testing against a Biotek Index 2 simulator and Masimo's simulator with signal strengths of greater than 0.02% and transmission of greater than 5% for saturations ranging from 70% to 100%.

WIRELESS TECHNOLOGY INFORMATION

Communication (Bluetooth)			
Туре	Bluetooth LE 5.0		
Frequency	2402-2480 MHz		
Classification of Output Power Rating	Conducted		
Max. Peak Output Power	6.13 dBm		
Output Power Type	Fixed at the Factory		
Modulation Types	GFSK		
Modulation Signals	Analog and Digital		
Available Data Rates	1 Mbps		
Recommended Max. Range	100 ft (~30 meters) line-of-sight		
Communication (Wi-Fi)			
Туре	WLAN Radio: IEEE 802.11 b/g/n		
Frequency	802.11b/g/n(HT20): 2412-2462 MHz 802.11n(HT40): 2422-2452 MHz		
Classification of Output Power Rating	Conducted		
Max. Peak Output Power	20 dBm		
Output Power Type	Fixed at the Factory		

Modulation Types	802.11b: DSSS 802.11g/n(HT20/HT40): OFDM		
Modulation Signals	Analog and Digital		
Available Data Rates	802.11b – 1, 2, 5.5, 11 Mbps. 802.11g – 6, 9, 12, 18, 24, 36, 48, 54 Mbps 802.11n- MCS0 – MCS7		
Security and Authentication			
Encryption	64/128-bit WEP, Dynamic WEP, WPA-TKIP, WPA2-AES		
Authentication	Open System, Shared Key, Pre-Shared Key (PSK), 802.1X: LEAP, PEAP, TTLS, TLS, EAP-FAST		
Radio Compliance			
USA	FCC ID: VFK-STORK		
Canada	IC ID: 7362A-STORK		

WARNING: Changes or modifications not expressly approved by the party responsible for compliance could void the user's authority to operate the equipment.

WARNING: The frequency bands of this device (2.4 GHz) are only for indoor use, in accordance with international telecommunication requirements.

GUIDANCE AND MANUFACTURER'S DECLARATION- ELECTROMAGNETIC EMISSIONS

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

EMISSION TEST	COMPLIANCE	ELECTROMAGNETIC ENVIRONMENT – GUIDANCE
RF Emissions (Radiated) CISPR 11	Group 1 Class B	The ME Equipment must emit electromagnetic energy in order t o perform its intended function. Nearby electronic equipment may be affected.
RF Emissions (Conducted) CISPR 11	Group 1 Class B	
Harmonic Emissions IEC 6 1000-3-2	Class A	Suitable for use in all establishments, including domestic environments and those directly connected to the public low-vol tage power supply network that supplies buildings used for dom estic purposes.
Voltage fluctuations/ Flicke r emissions IEC 61000-3-3	Complies	

GUIDANCE AND MANUFACTURER'S DECLARATION – ELECTROMAGNETIC IMMUNITY

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

IMMUNITY TEST	IEC 60601 TEST	COMPLIANCE L	ELECTROMAGNETIC ENVIRONMENT – GU
	LEVEL	EVEL	IDANCE
Electrostatic discharge (ESD) IEC 61000-4-2	+/- 8 kV contact +/- 15 kV air	+/- 8 kV contact +/- 15 kV air	Floors should be wood, concrete or ceramic ti le. If floors are covered with synthetic material , the relative humidity should be at least 30%.

Electrical fast transient / burst IEC 61000-4-4	+/- 1 kV for input/ output lines	+/- 1 kV for input/ output lines	Mains power quality should be that of a typical commercial or hospital environment.
Surge IEC 61000-4-5	+/-1 kV line(s) to li ne(s)	+/-1 kV line(s) to I ine(s)	Mains power quality should be that of a typical commercial or hospital environment
Conducted RF	3 Vrms	3 Vrms	Performed over 0.15-80 MHz
			Performed on the following ISM (industrial, sci entific and medical) bands of frequency: The bands between 0,15 MHz and 80 MHz are 6,7 65 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 amateur radio bands between 0,15 MHz and 80 MHz are 1,8 MHz
IEC 61000-4-6 6 '	6 Vrms		to 2,0 MHz, 3,5 MHz to 4,0 MHz, 5,3 MHz to 5
		6 Vrms	MHz, 7 MHz to 7,3 MHz, 10,1 MHz to 10,15 M Hz,
120 01000 4 0	O VIIIIS		14 MHz to 14,2 MHz, 18,07 MHz to 18,17 MH z, 21,0
			MHz to 21,4 MHz, 24,89 MHz to 24,99 MHz, 28,0
			MHz to 29,7 MHz and 50,0 MHz to 54,0 MHz
Power frequency (50			
/ 60 Hz) magnetic field	30 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of typical location in a typ
IEC 61000-4-8			ical hospital environment.

	0% UT1, 0.5	0% UT1, 0.5	
	cycle, at 0°, 45°,	cycle, at 0°, 45°,	
	90°, 135°, 180°,	90°, 135°, 180°,	
Valtaga dina an naugu	225°, 270°, and	225°, 270°, and	
Voltage dips on power supply input lines	315°;	315°;	
IEC 61000-4-11	0% UT 1 cycle,	0% UT 1 cycle,	
	and 70% UT 25/3 0 cycles at 0°	and 70% UT 25/3 0 cycles at 0	
			Mains power quality should be that of a
Voltage Interruptions on power supply input lines	0% UT, 250/300	0% UT, 250/300	typical commercial or hospital environment.
IEC 61000-4-11	cycle	cycle	
120 01000 4 11	1,7	1 - 3 - 1 -	
Radiated RF IEC 6100 0-4-3	10 V/m	10 V/m	Performed over 80 MHz to 2.7 GHz

Note 1: At 80 MHz and 800 MHz, 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. 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 acc uracy. 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 ME Equipment is used exceeds the applicable RF compliance level above, the ME Equipment should be observed to verify normal operation. If abnormal performance is observed, additional measures may be necessary, such as re-orienting or relocating the ME Equipment.

1 UT: Rated voltage for the equipment

ENCLOSURE PORT IMMUNITY TO RF WIRELESS COMMUNICATION EQUIPMENT

TEST FRE QUENCY	BAND (A) (MHZ)	SERVICE (A)	MODULATION (B)	MAXIMUM POWER (W)	DISTANCE (M)	IMMUNITY T EST LEVEL (V/M)
385	380-395	TETRA 400	Pulse modulation (b) 18 Hz	1.8	0.3	27
450	430-470	GMRS 460, FRS 46 0	FM (c) +/- 5 kHz deviatio n 1 kHz sine	2	0.3	28
710 745 780	704-787	LTE Band 13, 17	Pulse modulation (b) 217 Hz	0.2	0.3	9
810 870 930	800-960	GSM 800/900, TET RA 800, iDEN 820, CDMA 850, LTE Band 5	Pulse modulation (b) 18 Hz	2	0.3	28
1720 1845 1970	1700-1990	GSM 1800; CDMA 1900; GSM 1900; DECT; LTE Band 1, 3. 4. 35: UMTS	Pulse modulation (b) 217 Hz	2	0.3	28
2450	2400-2570	Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse modulation (b) 217 Hz	0.2	0.3	9

Note: If necessary to achieve the IMMUNITY TEST LEVEL, the distance between the transmitting antenna and the ME EQUIPMENT or ME SYSTEM may be reduced to 1 m. The 1 m test distance is permitted by IEC 61000-4-3.

- (a) For some services, only the uplink frequencies are included.
- (b) The carrier shall be modulated use a 50% duty cycle square wave signal.
- (c) As an alternative to FM modulation, 50% pulse modulation at 18 Hz may be used because while it does not r epresent actual modulation, it would be worst case.

RECOMMENDED SEPARATION DISTANCE BETWEEN PORTABLE AND MOBILE RF COMMUNICATION EQUIPMENT AND THE ME EQUIPMENT

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

RATED MAXIMUM OUTPUT POWER OF TRANSMITTER (W)

· · · · · · · · · · · · · · · · · · ·		
	d = 0.6*Sqrt (P)	
0.01	0.06	
0.1	0.19	
1	0.6	
10	1.9	
100	6	

For transmitters rated at a 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 m aximum output power rating of the transmitter in watts (W) according to the transmitter manufacturer..

Note 1: At 80 MHz and 800 MHz, 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.

The following symbols may appear on the product or product labeling:

Patents: http://www.masimo.com/patents.htm Stork is a trademark of Masimo Corporation. Masimo Corporation. Masimo Corporation. Masimo Corporation 52 Discovery Irvine, CA 92618 USA www.masimostork.com 0 2023 Masimo Corporation 304175/11839A-0623

Documents / Resources



MASiMO Stork Replacement Sensor [pdf] User Manual Stork Replacement Sensor, Replacement Sensor, Sensor

Manuals+,