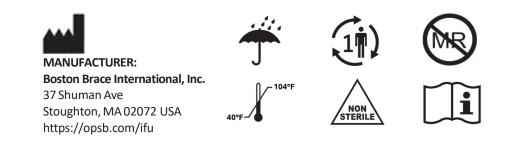




OPSB[™] Sensor System Instructions For Use







DESCRIPTION

The OrthoPediatrics Specialty Bracing (OPSB) Sensor System is intended to collect and transmit temperature data to aid patients and healthcare professionals (HCP) in their monitoring of a patient's compliance with their HCP's recommended brace or orthosis wear times and pattern. The system consists of the OPSB[™] Sensor, software applications (installable on Android and iOS phones), and cloud-based portal. The OPSB Sensor is a wireless accessory used to measure, collect and transmit measured temperature and time data of the Sensor. The OPSB[™] Sensor System is intended to be used in a clinical setting as well as outside of the clinical environment.

The OPSB sensor communicates, via Bluetooth, to an OPSB App available for Android and iOS platforms. Each OPSB Sensor has a unique QR code that allows secure synchronization to a corresponding smart phone containing the downloaded OPSB App. Synchronized data is automatically uploaded and stored to a cloud platform for analysis and display to the corresponding smart phone as a visual reward using stars and trophies for reaching wear goals. The cloud platform also allows authorized members of the patient's HCP team, via a secure log in, to remotely monitor the patient's wear times and patterns.

MATERIALS

The OPSB Sensor System is made from Polycarbonate/Acrylonitrile Butadiene Styrene Cycoloy C6600 (Sensor enclosure), FLEXCON 1400U (Sensor label), medical grade polyurethane (clear adhesive Sensor securing mechanism), and includes 3V Lithium coin-cell battery. There are no additional cables, transducers, or accessories.

INDICATIONS FOR USE

The OPSB Sensor System is intended to collect, transmit and store temperature data to monitor a patient's wear time and pattern.

CLIINICAL BENEFITS

The OPSB Sensor System provides valuable information regarding compliance with a HCP's recommended brace or orthosis wear time and pattern. Studies on brace and orthosis wear time show that when patients know they are being monitored and they receive a performance report, their adherence to the recommended hours of wear time improves. Achieving the goal is a time to celebrate. If there are some difficulties wearing the brace, then conversations between patient and HCP can be had to help patients meet wear time goals.

CONTRAINDICATIONS

Do not use the OPSB Sensor System in the presence of any contraindication. Contraindications include but are not limited to:

- Material sensitivity documented or suspected.
- Inability to follow treatment protocol and care instructions.

GENERAL SAFETY PRECAUTIONS

- Danger of damage to the OPSB Sensor: Do not attempt to modify or repair the Sensor. This may result in hazardous situations. Do not open the Sensor enclosure.
- Do not use the OPSB Sensor if it shows visible signs of damage, including damage to the QR Code.
- Protect the OPSB Sensor from mechanical shock. Keep the sensor away from; (1) aggressive chemicals or flammable substances, (2) heat sources like radiant heaters or fireplaces, (3) pets or pests, (4) babies and (5) chlorinated water, salt water, sunscreen and DEET based insect repellents.
- Do not use the sensor while bathing, swimming, surfing, or in a hot tub or sauna.
- Please contact Boston Brace International, Inc. for all repairs and modifications to the OPSB Sensor. Any unauthorized repairs or modifications to the OPSB Sensor may void the warranty and compliance of the equipment. Changes or modifications made to this equipment not expressly approved by Boston Brace International, Inc., may void the FCC authorization to operate this equipment.





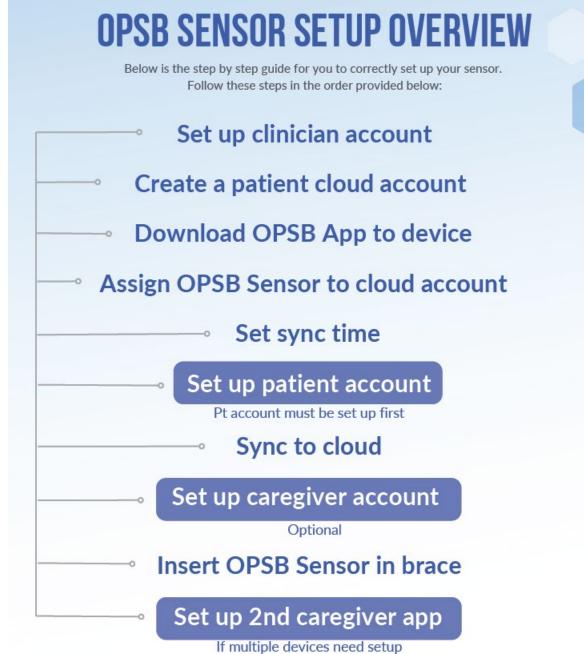
DEVICE SPECIFIC WARNINGS

- Do not attach the OPSB Sensor directly to the body, or to other parts of the brace not intended for Sensor attachment.
- To prevent skin irritation or direct skin contact, always wear a clean, dry garment.
- Harsh cleaners, such as bleach, are not recommended as the OPSB Sensor is in close contact with the skin while installed in the brace.
- Do not submerge the OPSB Sensor in water or cleaning liquids.
- Do not put the OPSB Sensor into a mechanical washer or dryer.
- Avoid misusing the OPSB Sensor as damage to the Sensor may cause battery fluid leak, which may lead to chemical burn. Do not touch a battery leakage. If you are exposed to battery leakage, wash the affected area thoroughly with clean water.
- Avoid potentially damaging circumstances: (1) do not drop the device, (2) do not damage the device, (3) do not connect to other devices, (4) do not put the device in contact with sharp objects.
- Avoid danger of electromagnetic interference: Do not use the OPSB Sensor close to MRI or CT facilities. Do not use the OPSB Sensor closer than 30 cm to a short wave, microwave or high frequency surgical device.
- Signal interference may be experienced if sensor is used within 30cm of another wireless device. See FCC Compliance Statement for more information.
- Sensor contains small parts, which could present a choking hazard.
- Apps (Android / iOS) are downloaded to the user's personal property and any vulnerabilities with the mobile devices are the responsibility of the user.
- App (Android / iOS) downloads must be made by and supervised by an adult, such as the patient's guardian where applicable.
- Loss of active Wi-Fi or cellular connection could impact data transfer.
- Patient data available through the Sensor, App and cloud platform can include name, date of birth, brace type, body temperature, and wear times. Boston Brace International, Inc. will keep confidential any patient identifying information that comes into its possession. Boston Brace International, Inc. may share and utilize limited or de-identified data with clinical partners of Boston Brace International, Inc. in research and in reporting outcome data to objectively review the impact of brace monitoring. Information regarding Boston Brace International, Inc.'s Privacy Policy may be found at: https://www.bostonoandp.com/privacy-policy/
- Patients may request deletion of their data from the cloud platform at any time by submitting a request to Boston Brace International Inc. through the Contact Us page: https://www.bostonoandp.com/contact/
- Patient data is stored in a third-party cloud platform. Data breaches, including cybersecurity attacks, can cause loss, damage, or unauthorized disclosure of patient data and impact the availability and accuracy of patient data. Patients and HCP are responsible for maintaining alternative sources and copies of patient data
- User credentials, including the names and contact information of patients, guardians, HCP, and authorized researchers are stored in the third-party cloud platform. Data breaches, including cybersecurity attacks, can cause loss, damage, or unauthorized disclosure of user credentials.





GUIDE FOR SETTING UP THE OPSB SENSOR SYSTEM







Patient registration by clinician/admin



Adding Sensor to patient account by clinician/admin







Patient Onboarding by clinician/admin







NOTE: Recommendation is to have Patient sign up on their own phone. If Patient doesn't have a phone, then Caregiver should download the OPSB app & sign in as Patient.

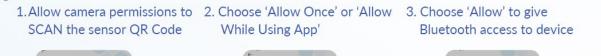
THE PATIENT APP ACCOUNT ORDER FOR THE SYN	
SETTING UP PATIENT	APP INSTRUCTIONS
1) Setting Up APP on a Patient Device	
1. Download the OPSB App off the App Store or Google Play store	OPSB
	OPSB Sensor



Instructions For Use OPSB[™] Sensor



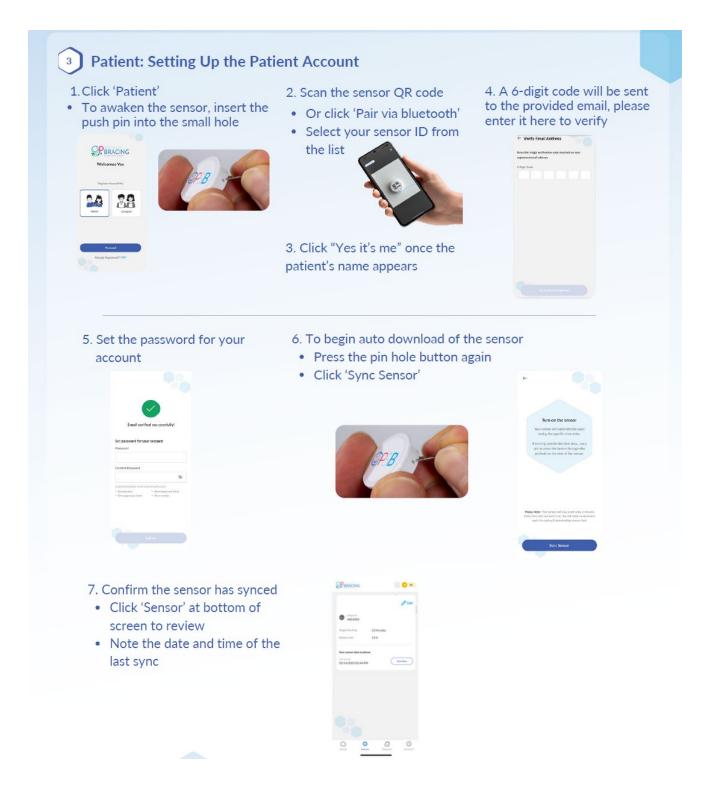
² Allowing APP Permissions





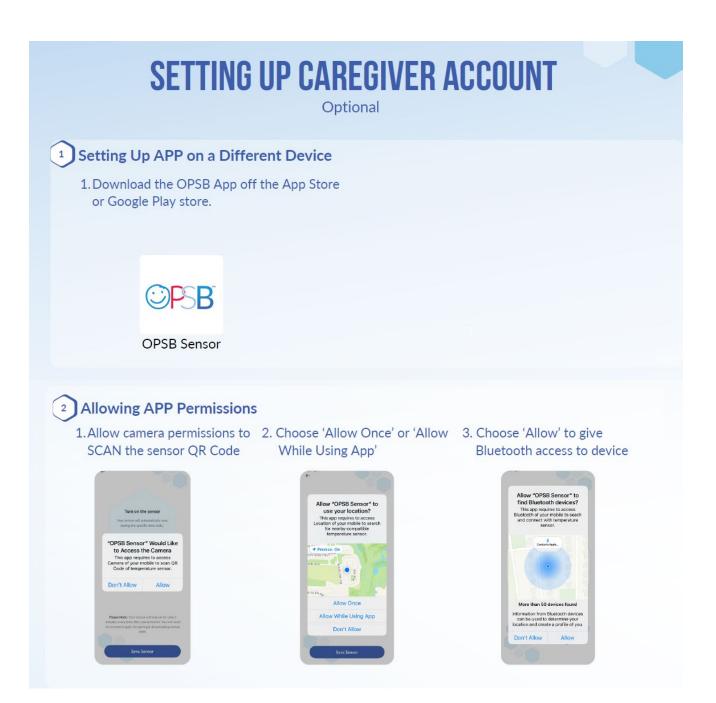






















Installing sensor into brace or orthosis

1 Insert the OPSB Sensor into the Orthosis

1. If the orthosis includes a pre-drilled hole then install, the OPSB Sensor by placing the sensor into the opening



2. If the orthosis does not include a predrilled hole, drill a 21mm hole using the appropriate drill bit. Then install the OPSB Sensor following the above directions.



² Cover the OPSB Sensor with a Clear Adhesive

Once the OPSB sensor is positioned in the orthosis, cover the sensor with clear adhesive. This will secure the sensor into the brace (sensor securing mechanism included).

• Send adhesive cover home with family if another caregiver's device is going to be setup



Peel off white backing in the center and position over sensor



Once placed, peel off remaining backing



Final sensor install

Additional Sensor System Information

1. Operating modes

- a. The OPSB Sensor will support two types of low power modes: (1) Power Down Mode where powering on ("wake-up") is allowed only by pin-hole button press, and (2) Sleep Mode, where wake-up is allowed by wake-up pin-hole button press or the pre-set sync schedule (as determined by your HCP).
- The OPSB Sensor will support two types of Bluetooth Low-Energy Modes: (1) BLE-discovery mode, and (2) BLE-off mode.
- c. The App will download data from the Sensor when the sensor is within 1-meter of the mobile device.

2. Performing standard functions

a. After successfully connecting the OPSB Sensor to the OPSB App on a mobile device, the user will be able to view their earned star points based on sensor wear time data and their actual wear time compared to their target wear time.





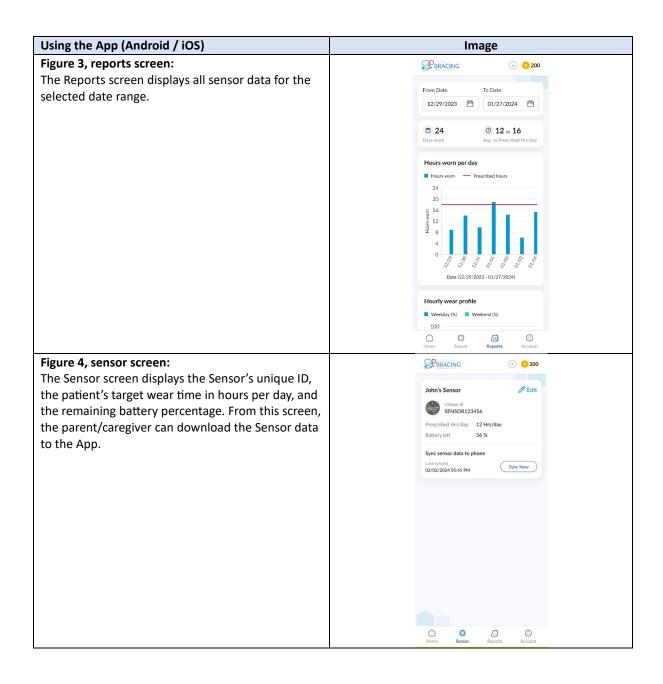
b. Additionally, after successfully connecting the OPSB Sensor to the OPSB App on a mobile device, the user can: (1) Navigate to the sensor tab or click the gray arrow to download sensor data to the APP, (2) Navigate to the reports screen to view all data and brace wear-trends, or (3) Navigate to the accounts tab to view or modify their account settings.

Using the App (Android / iOS)	Image
Figure 1, login screen: New users will proceed through the patient or caregiver workflow to create their account and connect with the Sensor. Existing users will proceed to the "login" screen and use their previously created credentials (email/password) to enter the App.	<image/> <image/> <image/> <section-header><section-header><section-header><section-header></section-header></section-header></section-header></section-header>
Figure 2, home screen: The Home screen displays the Sensor data, and all earned star points. Star points are earned once the patient reaches their target hours goal for the day. From this screen, the patient/caregiver can download the most recent Sensor data to the App. Background Data Syncing will occur automatically at the specific time slots set by the clinician, IF the APP remains open (in the background OR prior to the sync times)	Exercise Every John, Good Evening! 2000 Total star point Otal star points collected Hours worn per day Otal star points collected

3. Using the OPSB App (Android / iOS) -- Logging in after Registration











Using the App (Android / iOS)	Image
Figure 5, account screen: The Account screen displays the account details of the patient, and allows the patient to modify the App settings and to request deletion of their OPSB App account and all their stored data. Patients may request deletion of their data from the cloud platform at any time by submitting a request to Boston Brace International Inc. through the Contact Us page: <u>https://www.bostonoandp.com/contact/</u> .	7.04 Image: Second

4. DISPLAY INDICATORS AND ICONS

Display Indicator and icons	Image
The down-arrow and "Sync Now" buttons identified in the image on the right are used to initiate the data download process with the Sensor if syncing data outside of the scheduled sync window. For syncing outside of scheduled time slots: Awaken the sensor by pushing a pin in the small hole on the side of the sensor. Click "Sync Sensor"	Image: Sensor Image: Sensor Edit Image: Unique Id SENSOR123456 Prescribed Hrs/day 12 Hrs/day Battery left 56 % Sync sensor data to phone Last synced Q2/02/2024 05:45 PM Sync Now
	Home Sensor Reports Account

POSSIBLE ADVERSE EFFECTS

Patients should be advised to consult their HCP team immediately if experiencing any of the following:

- Skin irritation
- Pressure sores
- Pain, discomfort, or abnormal sensations due to the presence of the sensor

PATIENT COUNSELING

These Instructions for Use and all risks and benefits of the OPSB Sensor System based on individual patient factors should be fully explained to the patient by their HCP. Patient should contact their HCP in relation to the operation of the OPSB Sensor System. When necessary, the HCP may contact Boston Brace International, Inc. Please refer to the





Instructions for Use for the brace or orthosis being used with the OPSB Senor System for important information regarding the instructions, warnings, precautions, and possible adverse effects of the brace or orthosis.

MRI SAFETY INFORMATION

The OPSB Sensor System is magnetic resonance (MR) unsafe and should remain outside magnetic resonance imaging (MRI) scanner rooms.

STERILIZATION INFORMATION

The OPSB Sensor System cannot be sterilized via steam, gamma, ETO, or x-ray. The OPSB Sensor System is provided nonsterile and does not require sterilization before use.

CLEANING INSTRUCTIONS

Clean the outside of the OPSB Sensor with 70% rubbing alcohol. The Sensor should be allowed to air dry. Do not use a heat source for drying.

STORAGE AND HANDLING

Avoid exposing the OPSB Sensor to extreme temperatures. The ideal temperature range is 40°F to 104°F. Please see the above Precautions and Warnings for other storage and handling instructions.

REUSE / USE LIFE

All parts of the OPSB Sensor were designed to be used by a single patient, multi-use under normal use conditions and as instructed by a clinical care provider. The OPSB Sensor has a shelf life of two (2) years. After first powering on, the OPSB Sensor has a useful life of one (1) year.

SAFE DISPOSAL

When the OPSB Sensor reaches the end of useful life, recycle or dispose of the equipment according to appropriate local and regional regulations. Do not dispose of the Sensor in household refuse.

IMPORTANT STATEMENT

- The OPSB Sensor System manufacturer and distributor are not liable for cases of material damage or personal injury caused by misuse of the Sensor or non-compliance with these Instructions for Use. Normal use is defined as a single user following intended use.
- Instructions for Use (IFU) are available in English in paper copy as well as at https://opsb.com/ifu. To obtain additional copies of the paper IFU, please call Customer Service Group at Boston Brace International, Inc. at 1-800-262-2235.
- If a patient, user, or HCP becomes aware of a malfunction of the OPSB Sensor System or any component of the OPSB Sensor System, please immediately report such event to Boston Brace International, Inc.

CUSTOMER SUPPORT

 Contact Us at <u>customerservice@bostonoandp.com</u> Telephone: 1-800-262-2235





SYMBOL LEGEND

Symbol	Meaning	Symbol	Meaning
	Manufacturer	$\overline{\mathbf{x}}$	Date of manufacture
REF	Catalog number	SN	Serial number
	Single Patient, Multiple Use	NON STERILE	Non-sterile
Ĩ	Consult Instructions for Use or Consult Electronic Instructions for Use	MAR	MR Unsafe
40°F-	Temperature limit	MD	Medical Device

SENSOR LABELING

The following label is found on the rear side of the OPSB Sensor Packaging. The label measures 3 inches by 2 inches.



SAFETY

- 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.
- The sensor was successfully tested for compliance with the following tests: Radiated Emission, Electrostatic Discharge, Immunity to Radio Frequency Electromagnetic Fields, Proximity fields from RF wireless communications equipment, and Power Frequency and Magnetic field Test.
- There are no deviations from any standards used.
- No maintenance is needed to ensure basic safety and essential performance.

BATTERY RATING

• 3.0v dc, 130mA, 3W





FCC COMPLIANCE STATEMENT

FCC ID: 2BNBR-OPSB1. Changes or modifications to this product not expressly approved by the party responsible for compliance could void the electromagnetic compatibility (EMC) and wireless compliance and void the user's authority to operate the equipment. This device complies with Part 15 of the FCC Rules. Operation is subject to the following conditions: (1) this device may not cause harmful interference, (2) this device must accept any interference received, including interference that may cause undesired operation.

This equipment has been tested and found to comply with the limits for Class A digital device, pursuant to Part 15 of the FCC Rules. These limits are designed to provide reasonable protection against harmful interference in a residential installation. This equipment generates, uses and can radiate radio frequency energy and, if not installed and used in accordance with the instructions, may cause harmful interference to radio communications. However, there is no guarantee that interference will not occur in a particular installation.

If this equipment does cause harmful interference to radio or television reception, which can be determined by turning the equipment off and on, the user is encouraged to try to correct the interference by one or more of the following measures: (1) reorient or relocate the receiving antenna, (2) increase the separation between the equipment and receiver, (3) connect the equipment into an outlet on a circuit different to that to which the receiver is connected, and (4) consult the dealer or an experienced radio/TV technician for help.

Compliance Standards

Guidance and Manufacturer's Declarations – Electromagnetic Emissions		
IEC 60601-1-2:2014+A1:2020		
Emission Test	Home Healthcare Environment	
Radiated Emission	CISPR 11, Class B, Group 1	
Guidance and Manufacturer's Declarations – Electromagnetic Immunity		
Immunity Tests	Home Healthcare Environment	
Electrostatic Discharges (IEC 61000-4-2)	Contact Discharge: ±8kV	
	Air Discharge: ±15kV	
RF electromagnetic fields (IEC 61000-4-3)	80MHz-2700MHz, 10V/m, 80%, 1kHz	
Proximity fields from RF wireless communications equipment (IEC 61000-4-3)	Refer Note	
power frequency magnetic fields (IEC 61000-4-8)	30 A/m	

Test Specifications for Enclosure Port Immunity to RF Wireless Communications Equipment

Test Frequency	Band	Service	Modulation	Maximum Power	Distance	Immunity Test Level
(MHz)	(MHz)			(W)	(m)	(V/m)
385	380 - 390	TETRA 400	Pulse Modulation 18 Hz	1.8	0.3	27
450	430 – 470	GMRS 460, FRs 460	FM ±5 kHz deviation 1kHz sine	2	0.3	28
710			Pulse Modulation			
745	704 – 787	LTE Band 13, 17	217 Hz	0.2	0.3	9
780			217 112			
810		GSM 800/900,				
870	800 – 960	TETRA 800, IDEN 820, CDMA 850,	Pulse Modulation 18Hz	2	0.3	28
930		LTE Band 5				
1720		Bluetooth, WLAN,	Pulse Modulation			
1845	1700 – 1990	802.11 b/g/n, LTE	217 Hz	2	0.3	28
1970		Band 7	217 112			
2450	2400 – 2570	Bluetooth, WLAN, 802.11 b/g/n, RFID 2450, LTE Band 7	Pulse Modulation 217 Hz	2	0.3	28
5240	5100 - 5800	WLAN 802.11 a/n	Pulse Modulation	0.2	0.3	9
5500	5100 5800		217 Hz	0.2	0.5	5





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