

CORE 500™

Instructions For Use

CORE 500™
Digital Stethoscope



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1. General Information

This manual provides information to guide trained medical professionals in the safe and effective operation and proper maintenance of the CORE 500™ Digital Stethoscope. It's important that you read and understand all instructions in this manual before operating the device, and pay careful attention to the warnings and cautions throughout the manual.

Operate and maintain this product according to the safety and operating procedures in this manual, and only for its intended purpose. Always use the information in this document with sound clinical judgment and best clinical procedures.

Note: Depending on your platform, hardware, and country, certain features may not be available.

1.1 Indications and Intended Purpose

The CORE 500™ Digital Stethoscope is intended to be used by clinicians to electronically amplify, filter, and transfer body sounds and three lead electrocardiogram (ECG) waveforms. The CORE 500™ Digital Stethoscope also displays ECG waveforms and heart rate on the display and accompanying mobile application (when prescribed or used under the care of a clinician).

The data offered by the device is only significant when used in conjunction with clinician evaluation as well as consideration of other relevant patient data.

1.2 Device Description

CORE 500™ Digital Stethoscope (CORE 500™) is an electronic stethoscope with integrated electrodes for electrocardiogram (ECG). The device consists of a chestpiece, detachable earpieces (Eko Earpiece) and a mobile application (Eko App) and is intended as a digital auscultation tool on patients requiring physical assessment by the health care providers. CORE 500™ provides the ability to amplify, filter, and transfer body sounds with the chestpiece diaphragm, and three lead ECG through electrodes integrated around the chestpiece.

CORE 500™ features three auscultation modes for better auscultation experience by filtering acoustic data and enhancing the primary frequency range of particular body sounds: Cardiac Mode for heart sounds, Pulmonary Mode for lung sounds, and Wide Band Mode for general auscultation. CORE 500™ also detects and computes the heart rate in real-time based on the phonocardiogram (PCG) data. The computed heart rate and the ECG waveforms can be displayed on the screen mounted on top of the chestpiece, as well as the accompanying mobile application.

1.3 Clinical Benefit

The CORE 500™ Digital Stethoscope is a digital auscultation tool that improves the physical assessment of patients by health care providers.

As an integral part of a physical assessment, clinicians' interpretations of body sounds via the CORE 500™ Digital Stethoscope can help them rule in or out different pathological conditions in a patient. The integrated ECG lets the clinician quickly evaluate the ECG of the patient.

By helping physicians accurately detect the presence of conditions that warrant more investigation, further workup and testing can be better focused and more likely to result in the clinical benefit of an accurate diagnosis for the patient.

1.4 Continuous Operating Conditions

The operating range of the CORE 500™ is:

- A temperature range of + 5°C to + 45 °C.
- A relative humidity range of 15% to 90%, non-condensing, but not requiring a water vapor partial pressure greater than 50hPa.

It's recommended to avoid exposure to extreme heat, cold, solvents, and oils. Extreme heat and cold will negatively affect the lithium ion battery in the device and may affect battery life.

NOTE: If the device is used for a prolonged time at maximum ambient temperature it can get hot.

*(*CORE 500™ had an increase in temperature up to 47°C when tested at 45°C ambient temperature)*

1.5 Environmental Conditions of Transport and Storage Between Uses

The device is expected to be stored in a room with the following parameters:

- A temperature range of - 25 °C to + 5 °C.
- A temperature range of + 5 °C to + 35 °C at a relative humidity up to 90%, non-condensing.
- A temperature range of > 35 °C to 70 °C at a water vapor pressure up to 50 hPa.

1.6 System Requirements

For full functionality, the system requires users to connect their CORE 500™ with an internet-enabled smart mobile device using the Eko App. The app supports Apple® mobile devices. Make sure your system and mobile device meets or exceeds the minimum performance specifications (refer to Section 2.7 Technical Specification). Additional information on the most up-to-date system requirements can be found at support.ekohealth.com.

NOTICE: Some of the features of the Eko App require a minimum internet connection speed. The minimum recommended upload speed for the mobile app is 4000 Kbps. A minimum of 4G cellular data service or similar is recommended for the app. The app can be used to visualize waveforms and tracings without an internet connection, however an internet connection is necessary to save the data.

CORE 500™ uses Bluetooth® LE. Mobile devices used must be compatible with Bluetooth® LE.

Apple® is a registered trademark of Apple, Inc. Bluetooth® is a registered trademark of Bluetooth SIG, Inc.

1.7 Help and Assistance

For general and product-related comments, questions, or concerns, please contact Eko Devices, Inc., directly. If you have any questions or concerns about results found with the device, please consult a physician.

Serious Incident Reporting

If a serious incident has occurred in relation to the device, it should be reported to the manufacturer and the local competent authority in which the user and/or patient is established. A serious incident means any incident that directly or indirectly led, might have led, or, in case of recurrence, could lead to any of the following: the death of a patient, user or other person, the temporary or permanent serious deterioration of a patient's, user's, fetus's, or other person's state of health, or a serious public health threat.

Manufacturer Information

Eko Devices, Inc.
1212 Broadway, Suite 100
Oakland, CA 94612 USA

General Assistance and FAQs

support.ekohealth.com
Phone Support
(USA) 1.844.356.3384

Warranty Information

Eko provides a limited warranty for CORE 500™. Please visit ekohealth.com/warranty for a full description of the warranty.

Product Reference and Information

ekohealth.com

1.8 EMC Compliance**FCC Intentional Radiator Certification**

CORE 500™ Digital Stethoscope
FCC ID: 2ANB3-E8 IC: 23063-E8
MIC ID: PENDING CERTIFICATION

This equipment is approved by the FCC under the FCC ID numbers shown above. This device complies with Part 15 of the FCC rules. Operation is subject to the following two conditions: (1.) This device may not cause harmful interference, and (2.) This device must accept any interference received, including interference that may cause undesirable operation.

NO MODIFICATION: Modifications to this device shall not be made without the written consent of Eko Devices, Inc. Unauthorized modifications may void the authority granted under Federal Communications Commission rules permitting the operation of this device.

EMC Compliance Europe

This equipment complies with the EMC requirements of the IEC 60601-1-2.

FCC and Industry Canada Compliance Statement

This device complies with FCC Rules Part 15 and with Industry Canada license-exempt RSS standard(s). Operation is subject to two conditions: (1.) This device may not cause harmful interference, and (2.) This device must accept any interference that may be received or that may cause undesired operation.

Le présent appareil est conforme aux CNR d'Industrie Canada applicables aux appareils radio exempts de licence. L'exploitation est autorisée aux deux conditions suivantes : (1.) l'appareil ne doit pas produire de brouillage, et (2.) l'utilisateur de l'appareil doit accepter tout brouillage radioélectrique subi, même si le brouillage est susceptible d'en compromettre le fonctionnement.

2. Safety and Security

2.1 Symbols



Consult Instructions For Use



Consult Accompanying Documents



Manufacturer



Date of Manufacture and Country of Manufacture



Caution



Medical Device



Unique Device Identifier

IP44

IEC 60529 IP Rating 44
IP44 is protection Against ingress of solid foreign objects ≥ 1.0 mm diameter and splashing water



Authorized representative in the European Community/
European Union



MR Unsafe



CE Marking



WEEE separate collection facilities for recovery and recycling



Bluetooth Connectivity



Catalogue Number



Serial Number



Rx Only



Importer



Type CF Applied Part



Environmental conditions of transport and storage between
uses: -25 °C to $+70$ °C.



Environmental conditions of transport and storage between
uses: relative humidity up to 90 %, non-condensing

2.2 Precautions

- The device is intended to be used by or under the supervision of licensed medical professionals. The device may be used on patients during a physical assessment in a clinical setting or by patients under the supervision of a clinician. The system provides one source of data that is significant only when used in conjunction with clinician oversight and consideration of other relevant patient information. The ECG displayed on the device screen is a tool to assist clinical decisions and is not a diagnosis.
- CORE 500™ should be used only by qualified clinicians or by patients with an adequate understanding of the device. CORE 500™ is intended for use on patients that can be auscultated normally with an acoustic stethoscope.
- This manual provides instructions for the use of CORE 500™ and the Eko App. It's assumed that the user is familiar with basic mobile application use on iOS™ and Android devices.
- Standard procedures for auscultation should be followed, including background noise reduction and optimal patient positioning. Use the provided earpiece with the CORE 500™ for best audio quality. The quality of organ sounds is dependent on proper use, including holding the device still and increasing volume as needed.
- The quality of the ECG is dependent on proper preparation practices including, but not limited to, cleaning the contact area, electrodes and using ECG gel. If used on a portion of the body with significant body fat, body hair, or very dry skin, a successful recording may not be possible.
- The device display is not recommended to determine the absolute amplitude of the ECG signal. Eko recommends that the display be primarily used for ensuring good ECG signal quality. The accompanying mobile app should be used to read and interpret the three lead ECG.
- The device can be used with any wired headphones or wired headsets. No performance guarantees are claimed using other audio products. If using other headphones, insertable earbuds provide the best sound quality. The device can also be used with wireless listening devices, such as hearing aids, connected through the mobile app. For optimal audio quality while using the mobile app, it's not recommended to listen through the mobile device's in-built speaker.
- Please read, understand, and follow all safety information contained in these instructions prior to using the CORE 500™. It's recommended that these instructions be retained for future reference.
- No modification of this equipment is allowed. There are no repairable parts inside the CORE 500™.
- This device does not detect or measure all heart rate, heart rhythm, and heart waveform changes.
- DO NOT use the device while it is charging.
- To reduce the risks associated with infection, follow all cleaning instructions included in this manual. Establish and follow a cleaning schedule after each use.
- DO NOT use the device over broken skin or wound areas.
- DO NOT continue to use if you have an allergic reaction to the device materials or if your skin appears irritated or inflamed after use. Check with a healthcare professional before restarting use.
- To reduce the risks associated with inaccurate data acquisition, store and operate this device only as instructed in this manual.

- It's recommended that the battery be recharged within 30 minutes of the low battery indicator warning. Recharge the battery using only the appropriate USB-C cable.
- DO NOT immerse the device in a liquid or subject it to any sterilization processes other than those described in this manual. The device is non-sterile.
- To reduce the risk of device interference, keep the device at least two meters away from all RF emitters, including Wi-Fi routers and radios when operating or charging.
- To reduce the risks associated with very strong electromagnetic fields, avoid using the device near strong radio frequency (RF) signals or portable and/or mobile RF devices.
- If sudden or unexpected sounds are heard, move away from any radio transmitting antennas. Using accessories, transducers, and cables not produced by Eko may result in increased RF emissions or decreased immunity.
- The device contains a Bluetooth Class 2 wireless data link. The maximum radio frequency field strength generated by the device is below three volts per meter, a level that is considered safe to use with other medical devices. However, audio, video, and other similar equipment may cause electromagnetic interference. If such devices are encountered and cause interference, immediately move the device away from that device and/or turn the Bluetooth feature of the interfering device OFF.
- The device uses a Bluetooth Class 2 wireless data link. The Bluetooth range will be reduced when objects (walls, furniture, people, etc.) are between the device and a paired mobile device. To improve Bluetooth connection, reduce the distance and/or allow a line of sight between the device and the mobile device.
- To reduce the risks associated with environmental contamination, follow applicable regulations when disposing of this device. The device contains a rechargeable battery. Please properly dispose of the device as mandated by local directives.
- Disperse any static electricity before using the unit.
- Do not operate or store the CORE 500™ in extremely hot, cold, humid, or wet conditions.
- The ECG and body sounds should be used in conjunction with a clinical evaluation. Do not use as the sole basis for medication or treatment decisions.
- Never use the stethoscope without eartips firmly locked in place.
- CORE 500™ is not intended for use with flammable anesthetics or flammable agents.

2.3 Warnings

Failure to follow caution and warning could result in damage to the internal components of the device. Internal damage to the product could cause malfunction of the product, possibly leading to complete loss of function. If problems are encountered with the device, do not attempt to repair it. Please notify our support team for assistance.

- **WARNING:** Stethoscope tubing can be a strangulation hazard. Keep away from unsupervised children.
- **WARNING:** Eartips can be swallowed and cause a choking hazard. Ensure all parts are properly attached and stored.

- **WARNING: MR-unsafe!** Do not expose the device to a magnetic resonance (MR) environment. The device may present a risk of projectile injury due to the presence of ferromagnetic materials that can be attracted by the MR magnet core. Thermal injury and burns may occur due to the metal components of the device that can heat during MR scanning. The device may generate artifacts in the MR image. The device may not function properly due to the strong magnetic and radio frequency fields generated by the MR scanner.
- **WARNING:** The CORE 500™ is not intended to be used in an oxygen-rich environment.
- **WARNING:** DO NOT use adjacent to or stacked with other equipment because it could result in improper operation.
- **WARNING:** DO NOT use unapproved accessories. Use of non-Eko approved accessories or transducers and cables could result in electromagnetic emissions or decreased electromagnetic immunity of this device and result in improper operation.
- **WARNING:** DO NOT use portable RF communications equipment (including peripherals such as antenna cables and external antennas) closer than 30 cm (12 inches) to any part of the CORE 500™. Otherwise, degradation of the performance of the CORE 500™ could result.
- **WARNING:** Only connect to a power supply to mains voltage marked appropriately in it. During charging, the power supply needs to be easily accessible, in case there is a need for disconnection from mains in an unexpected incident.

2.4 Network Security

When connecting your smart device, use a network that supports Wi-Fi 802.11n. It is recommended to secure this network using WPA (Wi-Fi Protected Access) or WPA2 (Wi-Fi Protected Access II) as your security protocol. For information on setting up your wireless network security, refer to your network equipment's documentation.

All data transmitted from the Eko App is encrypted in transit using TLS 1.2 or greater, and all data is encrypted at rest using AES 256.

In addition to security features embedded in the system, it's highly recommended that users of the Eko App and Eko Dashboard use networking security features to protect patient data created and stored using this software. Common examples include strong passwords, biometric authorization, two-factor authentication, and VPN encryption when available.

2.5 Patient Privacy

The privacy of patient health information may be protected by state, federal, or international/foreign laws that regulate how such information can be used, stored, transmitted, and disclosed. The Eko system employs security features that are compliant with HIPAA policies. Third-party access may be prohibited to such information without obtaining written authorization from the patient.

The user is fully responsible for understanding and following all laws that regulate storage, storage transmission, and disclosure of any electronic patient data through the use of software. If the user becomes unable to comply with a law or restriction that applies to use and disclosure of such data, the user should not proceed to collect or save such information.

This application may require entry of individually identifiable health information in order to function. Records are stored and recalled through the use of patient name, date of birth, and/or patient ID number. By entering this information, the user assumes any and all risks of and liabilities incurred with using or transmitting such information.

2.6 Guidance and Manufacturer's Declaration – Electromagnetic Emission

The CORE 500™ is intended for use in the electromagnetic environment specified below. The user of the CORE 500™ should ensure that it is used in such an environment.

Applicable Emissions Test	Compliance	Electromagnetic Environment – Guidance
RF emissions CISPR 11	Group 1	CORE 500™ uses RF energy only for its internal functions. Therefore, its RF emissions are very low and are not likely to cause any interference in nearby electronic equipment. CORE 500™ is suitable for use in all establishments, including domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes.
RF emissions CISPR 11	Class B	
Harmonic Emissions IEC 61000-3-2	Not Applicable	
Voltage fluctuations/flicker emissions IEC 61000-3-3	Not Applicable	

Guidance and Manufacturer's Declaration – Electromagnetic Immunity

The CORE 500™ Digital Stethoscope is intended for use in the electromagnetic environment specified below. The user of the CORE 500™ Digital Stethoscope should ensure that it is used in such an environment.

Immunity Test	IEC 60601-1-2 Test Level	Compliance Level	Electromagnetic Environment – Guidance
Electrostatic Discharge (ESD) IEC 61000-4-2	+/- 8 kV contact +/- 15 kV	+/- 8 kV contact +/- 15 kV air	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/Burst IEC 61000-4-4	+/- 2 kV for supply lines +/- 1 kV for input/ output lines	Not Applicable	Not Applicable
Surge IEC 61000-4-5	+/- 1 kV line(s) to line(s) +/- 2 kV line(s) to earth	Not Applicable	Not Applicable
Voltage dips, short interruptions and voltage variations on power supply input lines IEC 61000-4-11	100% drop in UT for 0.5 cycle 0/45/ 90/135/ 180/225/ 270/315 degrees, 100% dip in UT for 1 cycle, 30% dip in UT for 25 cycle, 100% drop in UT for 5 sec	Not Applicable	Not Applicable
Power frequency (50/60 Hz) magnetic field IEC 61000-4-8	30 A/m	30 A/m	Power frequency magnetic fields should be at levels characteristic of a typical location in a typical commercial magnetic field or hospital environment.

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

The CORE 500™ Digital Stethoscope is intended for use in the electromagnetic environment in which radiated RF disturbances are controlled. The user of the CORE 500™ Digital Stethoscope can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the CORE 500™ Digital Stethoscope 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 2.5 GHz $d = 2.3 \sqrt{P}$
0.01	0.12	0.12	0.23
0.1	0.37	0.37	0.74
1	1.2	1.2	2.3
10	3.7	3.7	7.4
100	12	12	23

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 maximum power rating of the transmitter in watts (W) according to the transmitter manufacturer.

NOTE 1: At 80 MHz and 800 MHz, 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.

2.7 Technical Specifications

General Performance	
Audio Frequency Response	Bandwidth of 20 Hz - 2000 Hz
Audio Playback Volume	Output level from 85 to 100 dB SPL A-weighted with user selectable volume levels
User Volume Protection	Max Sound Pressure Level (SPL) output of 100 dB, SPL A weighted
ECG Performance	2 channel ECG in real-time in the 0.1 - 250 Hz frequency range
Heart Rate	+/- 5BPM mean absolute error
Expected Service Life	2 years for CORE 500™ system
Applied Parts	Type CF. The 3 electrodes and a left leg electrode on the bottom face of the device are type CF applied parts
Mode of Operation	Continuous

Bluetooth Characteristics	
Bluetooth Protocol	Support communication with supported Bluetooth Low Energy (BLE) 4.2 and BLE 5.0 clients
Data Transfer Encryption	Data transferred via Bluetooth is encrypted
Power	
Battery Type	Internally powered using rechargeable 3.7 V Lithium-ion polymer cell
Battery Life	<80% of battery capacity self drains in 6 months Minimum 5 hours continuous use
Physical Characteristics	
Dimensions	27 inches (685mm) long
Weight	6.6 ounces (186g) with earpiece
Environmental Specifications	
Environmental Conditions of Transport and Storage Between Uses	- 25 °C to + 5 °C + 5 °C to + 35 °C, relative humidity up to 90 %, non-condensing > 35 °C to 70 °C at a water vapor pressure up to 50 hPa (conforming to IEC 60601-1-11 and 60601-2-47)
Continuous Operating Conditions	5 °C to + 45 °C; relative humidity range of 15 % to 90 %, non-condensing (conforming to IEC 60601-1-11 and 60601-2-47)
Ingress Protection	IP Rating 44 IP44 is protection Against ingress of solid foreign objects \geq 1.0 mm diameter and splashing water
User Interface	
Chestpiece	Hand-held device with capacitive touch, mode button, volume button, and USB-C port (charging only). CORE 500™ can be safely charged* using a Certified USB-IF, Class II Double insulated USB charging port with output voltage rated at 4.75V-5.25V and charging current at 500mA - 2A. *(CORE 500™ was tested with an Apple Model:A1385 USB Power Adapter.)
Mobile Device	iPhone with iOS 6.1 and above
Earpiece	Standard 3.5 mm female TRS jack

3. Installing the Eko App

3.1 Downloading and Installing

The Eko App helps you to easily connect to CORE 500™ for secure transmission and analysis of your recordings.

Downloading the Eko App allows you to:

- Pair the CORE 500™ to your mobile device.
- Listen wirelessly through a headset.
- View PCG and three lead ECG visualizations.
- Start, save, and share recordings.
- Take advantage of additional usage guides.

After you install the Eko App, complete the CORE 500™ device setup on the app. The app walks you through setting up and using your CORE 500™.

Download the Eko App below:

Or, scan the QR code below to download Eko App:



3.2 Connecting CORE 500™ with App

1. Turn on your phone's Bluetooth.
2. Turn on your CORE 500™.
3. Open the Eko App and sign in.
4. Follow the onscreen instructions for pairing your device.
5. The Bluetooth icon appears on the CORE 500™ screen when connected.

4. Using the CORE 500™

4.1 Removing from Packaging

Carefully remove the device from the packaging. Before use, inspect the device for any damage. Do not use a damaged device.

There is no requirement to warm up the device prior to use.

4.2 Skin Preparation

Excessive hair, dirty skin, dry skin, or oily skin can impact the quality of the ECG tracing. Wetting the patient's skin with 70% isopropyl alcohol wipes can improve ECG electrode contact. Do not use the CORE 500™ over wound areas or areas of broken skin. Rub the skin vigorously to increase capillary blood flow to the tissues. ECG gels or saline solutions can also be used on the electrodes to improve signal quality.

4.3 Contents

This package includes:

- 1 CORE 500™ Digital Stethoscope
- 1 USB-C cable
- 1 Eko earpiece
- 4 silicone rubber ear tips (2 large, 2 small)
- 10 Alcohol wipes
- 1 Quick start guide
- 1 Instructions for Use

4.4 Earpiece Setup

Attach Earpiece

1. Plug the earpiece into the CORE 500™.
2. Twist the earpiece clockwise with a quarter turn until it locks.

Warning: Do not use excessive force when twisting the earpiece.

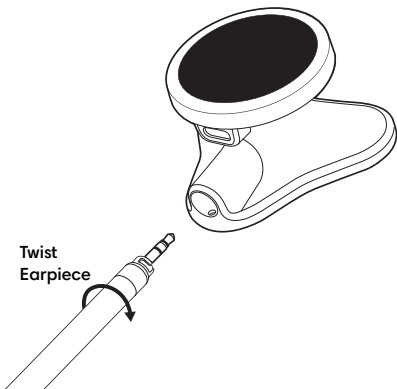
Detach Earpiece

1. Twist the earpiece counterclockwise with a quarter turn until it unlocks.
2. Remove the earpiece from the CORE 500™.

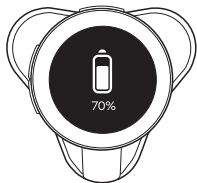
Warning: Do not use excessive force when twisting the earpiece.

Fit Earpiece

Use the right ear tip size. Try out the small or large ear tips for the best fit.



4.5 Charging



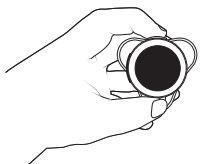
1. Connect the CORE 500™ to a power source using the included USB-C cable and a power adapter (not included).
2. The battery indicator shows the charge percentage.

The battery life is subject to use and is expected to last for five hours of continuous use. It takes approximately three hours to fully charge the device from 0%.

The CORE 500™ should be periodically recharged even when in storage. Lithium ion batteries slowly lose charge when in storage and may fall to an unacceptably low level, damaging the battery.

NOTE: The CORE 500™ will not operate or connect to the Eko App while charging.

4.6 Turning On and Off



Turn On

Pick up the device by placing your fingers around the space between the device face and electrodes. The CORE 500™ will turn on automatically.

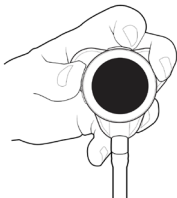
Or, press the top button to turn on the CORE 500™.

Sleep

The CORE 500 will automatically sleep within 15-60 seconds of inactivity. Pick it up to wake it up again. You can turn off this setting in the Eko App.

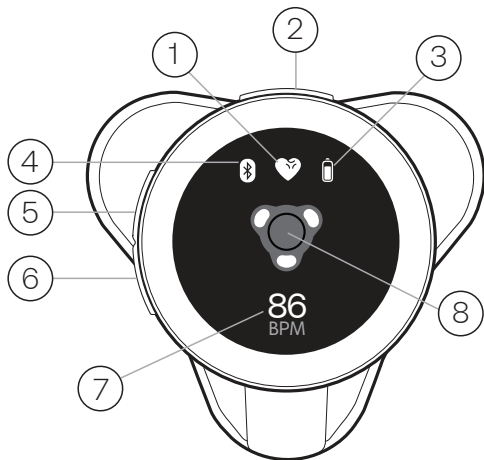
Turn off

Hold the top button while pressing the volume-up button once to turn it off.



Once you see the confirmation screen, press the volume-up button again to confirm.

Resting State



1. Current audio filter:

- Wide (for all use cases)
- Cardiac (with ECG)
- Pulmonary (with lung sounds)

2. Top button:

a. Start recording:

= press for 2 seconds

b. Change filter:

= press for <1 second

3. Battery level

4. Bluetooth connected

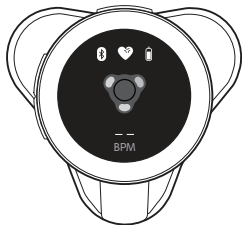
5. Volume up button

6. Volume down button

7. Real-time heart rate in beats per minute

8. Leads indicator:

- a. O = No skin contact detected
- b. ✓ = Skin contact detected

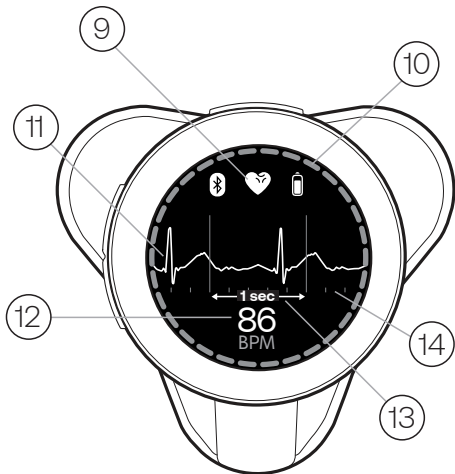


If the electrodes do not make contact with skin, the lead indicator will be gray.



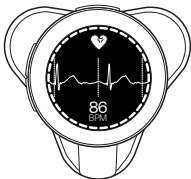
If the electrodes make contact with skin, the lead indicators will turn blue.

Active State



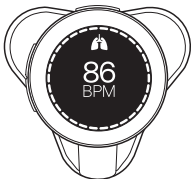
- 9. Current audio filter
- 10. Recording in progress
- 11. ECG waveform (shown in cardiac and wide modes)
- 12. Real-time heart rate in beats per minute
- 13. 1-second intervals
- 14. 0.2-second intervals

Each filter mode described below accentuates particular body sounds while auscultating: The cardiac mode is weighted toward heart sounds, pulmonary toward lung sounds, and wide band provides coverage for both.



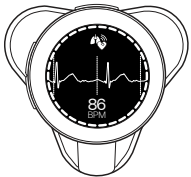
Cardiac audio filter

In the cardiac audio filter mode, the CORE 500™ screen shows a heart icon at the top, the ECG waveform for a two-second interval, and the real-time heart rate in beats per minute. If a recording is in progress, blue bars light up around the screen as the recording progresses.



Pulmonary audio filter

In the pulmonary audio filter mode, the CORE 500™ screen shows a lung icon at the top and the real-time heart rate in beats per minute. If a recording is in progress, blue bars light up around the screen as the recording progresses.

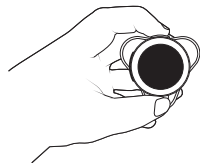


Wide audio filter

In the wide audio filter mode, the CORE 500™ screen shows an icon with a heart and lungs, the ECG waveform for a two-second interval, and the real-time heart rate in beats per minute. If a recording is in progress, blue bars light up around the screen as the recording progresses.

4.8 Changing the Volume

The device's sound level can be amplified in seven increments. Change the volume level by pressing the top (+) and bottom (-) of the volume button on the side of the device. The volume change is confirmed on the screen of the device.



4.9 Changing the Audio Filters

Press the top button for less than one second to switch to a different audio filter. You must press for less than one second between each audio filter. The available filters are cardiac (with ECG), pulmonary (with lung sounds), and wide (for all use cases) .

Example

The device is in cardiac mode. You press the top button for less than one second. The audio mode changes to pulmonary mode. You press the top button for less than one second. The audio mode changes to wide mode.

4.10 Capturing Sounds and ECGs

To capture sounds and ECGs, the CORE 500™ can be used on various locations and orientations of the chest. Each position will produce a unique body sound and ECG tracing. For ECG, place the device directly onto the patient's skin. Do not perform an ECG over the patient's clothing.

Audio

Capture sounds by placing the CORE 500™ anywhere on the body. For best audio, press the device firmly against the patient to reduce movement.

ECG

Capture the ECG signal by placing the CORE 500™ on the skin. The audio filter icon should be at the top, facing up. One position that works well is the left upper sternal border (next to the left sternum edge between the second and third rib), as shown in the placement illustration below. If the patient has particularly dry skin, significant body fat or chest hair, then alcohol wipes or conductive gel used with other ECG systems may be applied to CORE 500™ electrodes to improve the quality of the ECG signal.

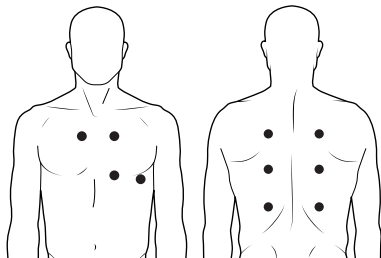
The device will confirm all electrodes have contact with the patient's skin by the three gray electrode icons turning blue simultaneously as they detect skin contact. If all three electrodes are gray, there is not sufficient skin contact.

Note that device display is not recommended to determine the absolute amplitude of the ECG signal. We recommend that the display be primarily used for ensuring good ECG signal quality. The accompanying mobile app should be used to read and interpret the three lead ECG.

Placement

CORE 500™ streams a 3-lead ECG. Best placement can vary. ECG electrodes must be placed on the skin. CORE 500™ can also be used to auscultate at all anatomical positions.

The dots (right) indicate generally accepted CORE 500™ placement positions. CORE 500™ is sensitive to vibration and hand movement. Remember to apply firm and constant pressure against the body to ensure good contact.



4.11 Starting a Recording

From the CORE 500™

Press the top button for two seconds to start a recording. The bars encircling the device interface light up blue to indicate the recording is in progress.

Note: To initiate a recording from the device, the device must be connected to the Eko App via bluetooth.

From the Eko App

Launch the app and click the Record button.

Note: To initiate a recording from the app, the device must be connected to the Eko App via bluetooth.

5. Processing, Cleaning, and Disposal

The CORE 500™ is a multiple patient, multiple use device.

Processing between uses

The CORE 500™ should be sufficiently charged and disinfected between uses following instructions provided in this manual. The CORE 500™ does not perform nor require periodic self-check maintenance activity to maintain functionality.

Ensure the CORE 500™ is within the specified Continuous operating conditions (section 2.7) prior to use.

There is no warm-up or cool-down period required between uses.

There are no known adverse effects of lint, dust, or direct sunlight on the functionality of CORE 500™. However, the device should be stored in a clean location.

The device should be stored away from children and pets to prevent unintended tampering.

CORE 500™ should not be reused if:

- The device enclosure or attachment has visible damage.
- The device does not turn ON/OFF.
- The device cannot be sufficiently charged.
- The device exhibits acoustic or ECG issues.
- The device exhibits other operational anomalies.
- The device packaging has visible damage, contamination, unintentional opening or exposure to environment conditions outside of specification.

Cleaning

All external surfaces of the hardware can be cleaned with isopropyl alcohol wipes. Under normal conditions, it is not necessary to remove the chestpiece from the earpiece during the cleaning procedure.

Ensure all external surfaces are dry prior to use.

NOTE: DO NOT immerse the device in any liquid or subject it to any high-pressure/autoclave sterilization processes.

Disposal

The device should not be discarded as unsorted waste but must be sent to separate collection facilities for recovery and recycling according to applicable local or national laws.

6. Manufacturing and Regulatory Information



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Made in Malaysia

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LBL390 Rev 2.0 - February 2023