

# OXXIOM™ USER MANUAL



**29826 Avenida de Las Banderas, Suite 300  
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# PACKAGE CONTENT

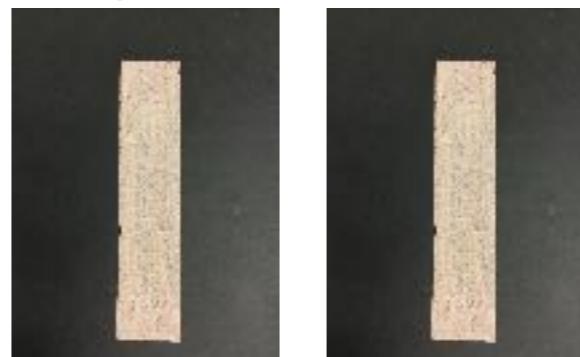
Each OXXIOM pack contains



① One User Manual



② Six 2 x 6 in self-adhering breathable gentle tapes



③ Two barcode packages with one OXXIOM unit each



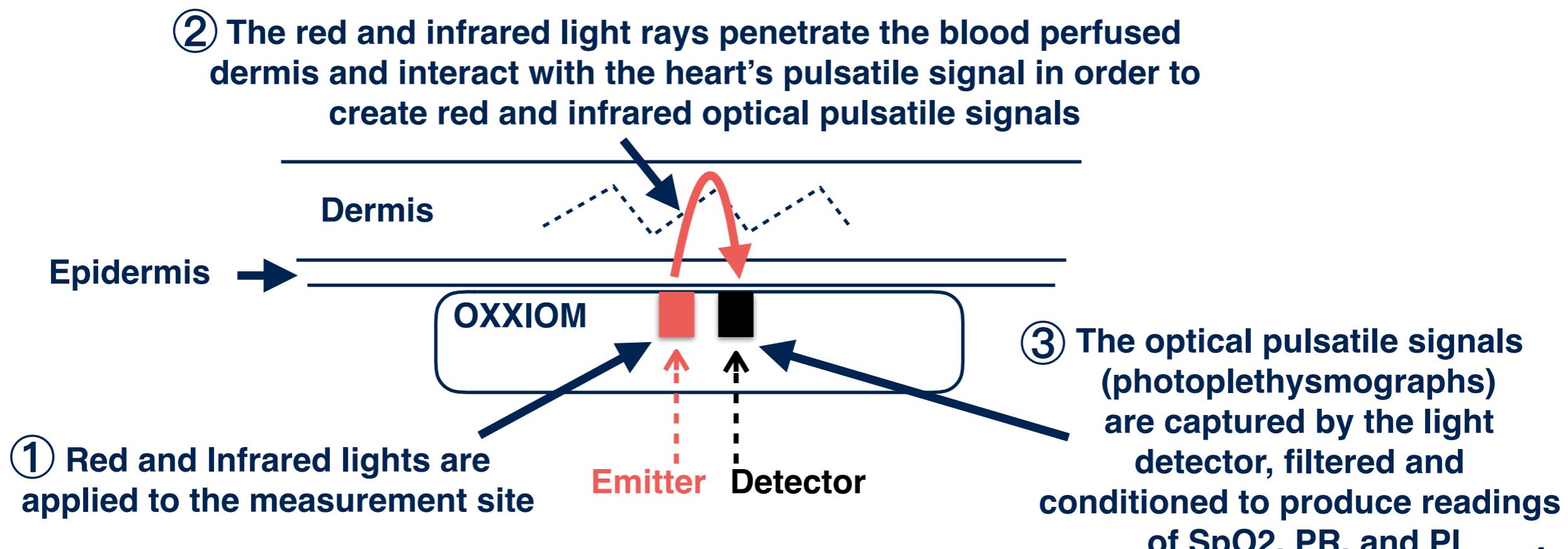
# OXXIOM OPERATION

Oxxiom's innovative design has combined the latest electronics and biosensing technology for the measurement of the patient's functional arterial oxygen saturation (SpO<sub>2</sub>), perfusion index (PI), and pulse rate (PR).

The device is a cableless, completely wireless, continuous, fully disposable, single-use pulse oximeter with the combined function of a clinical grade pulse oximeter monitor and reflective sensor into a single disposable device. Oxxiom is a small, lightweight, easy to use, battery operated pulse oximeter that offers 24 hours of continuous monitoring.

Oxxiom wirelessly transfers the patient's data to a host device (such as an iPhone or an iPad). The host device functions as a primary display and is required for Oxxiom's operation. Measurements are continuously taken by Oxxiom and the data is simultaneously and wirelessly transferred to the host device for display, alarms, and storage.

## Measurement Principle



# TECHNICAL SPECIFICATIONS

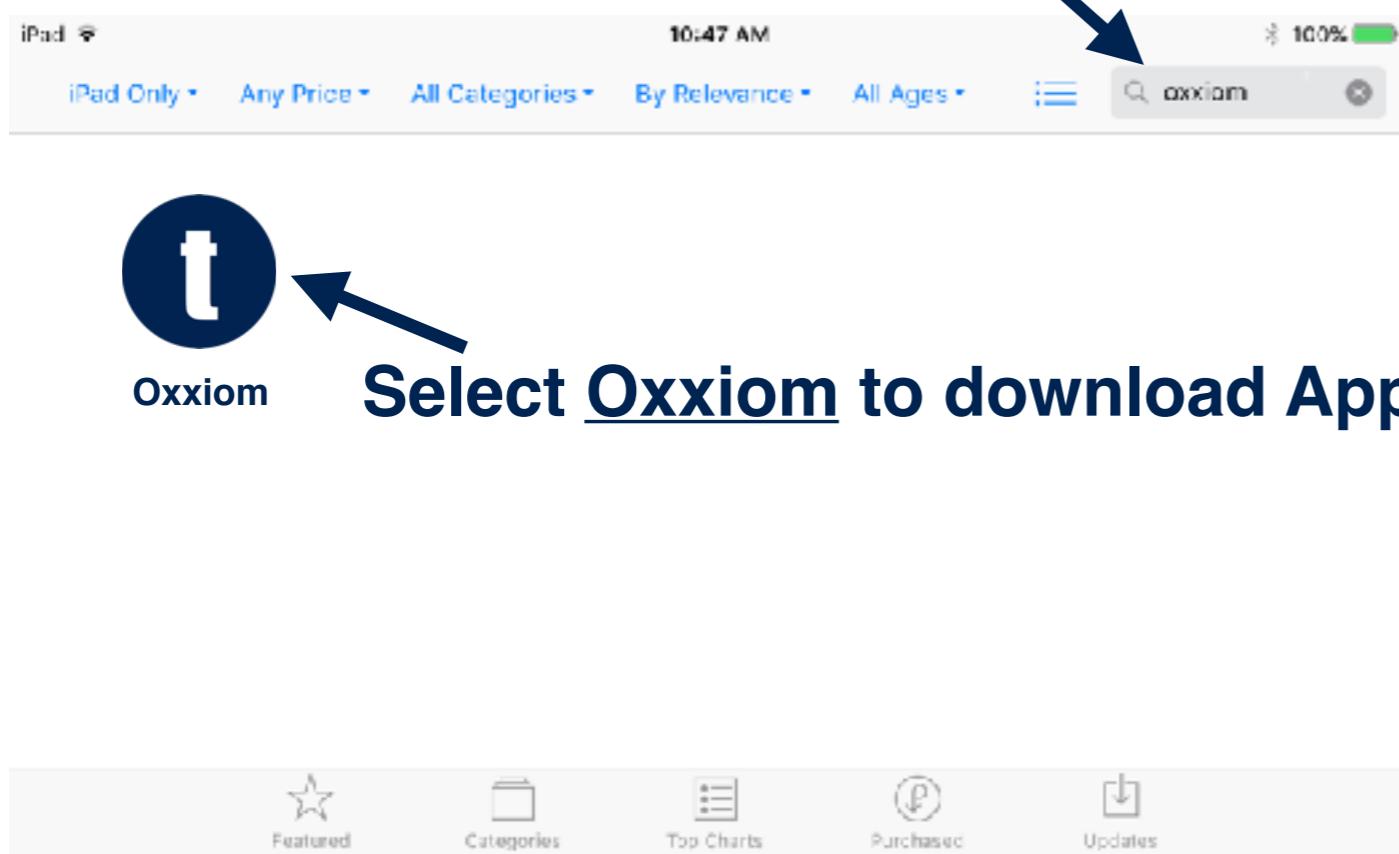
Features	Specifications
<b>Safety</b>	
Electrical, EMC, Pulse Oximetry, Biocompatibility	IEC 60601-1, IEC 60601-1-2, ISO 80601-2-61, ISO 0993-5, ISO 10993-10
<b>Physical</b>	
Dimensions, Weight	30 x 17 x 7.5 mm (12 x 0.7 x 0.3 in), 3.5 g (0.12 oz)
<b>Optical Sensor</b>	
Emitted light energy, Wavelength range	≤ 15 mW, 500-1,000 nm
<b>Accuracy per ISO 80601-2-61</b>	
SpO2 (70-100%)	+/- 3% on 68% of the population
PR (25-250 BPM)	+/- 3 BPM on 68% of the population
<b>Measurements Range on Host Device</b>	
SpO2	1-100%
PR	25-250 BPM
PI	0.03-20%
<b>Alarms on Host Device</b>	
SpO2, PR	Configurable high and low limits
<b>Environmental</b>	
Operating temperature, Operating humidity, Operating altitude	5-40°C (41-104°F), 95% RH max. at 40°C (104°F), Up to 3,012m (9,882 ft)
Storage temperature, Storage humidity, Storage altitude	0-60°C (32-140°F), 95% RH max. at 60°C (140°F), Up to 5,574m (18,228 ft)
<b>Wireless</b>	
Protocol, Range	Bluetooth Low Energy (BLE), 10 m (33 f)
<b>Battery</b>	
Technology, Nominal voltage, Nominal capacity	Non-rechargeable lithium/manganese dioxide (LiMnO2), 3 V, 150 mAh
Life	24 h of continuous monitoring

# WARNINGS

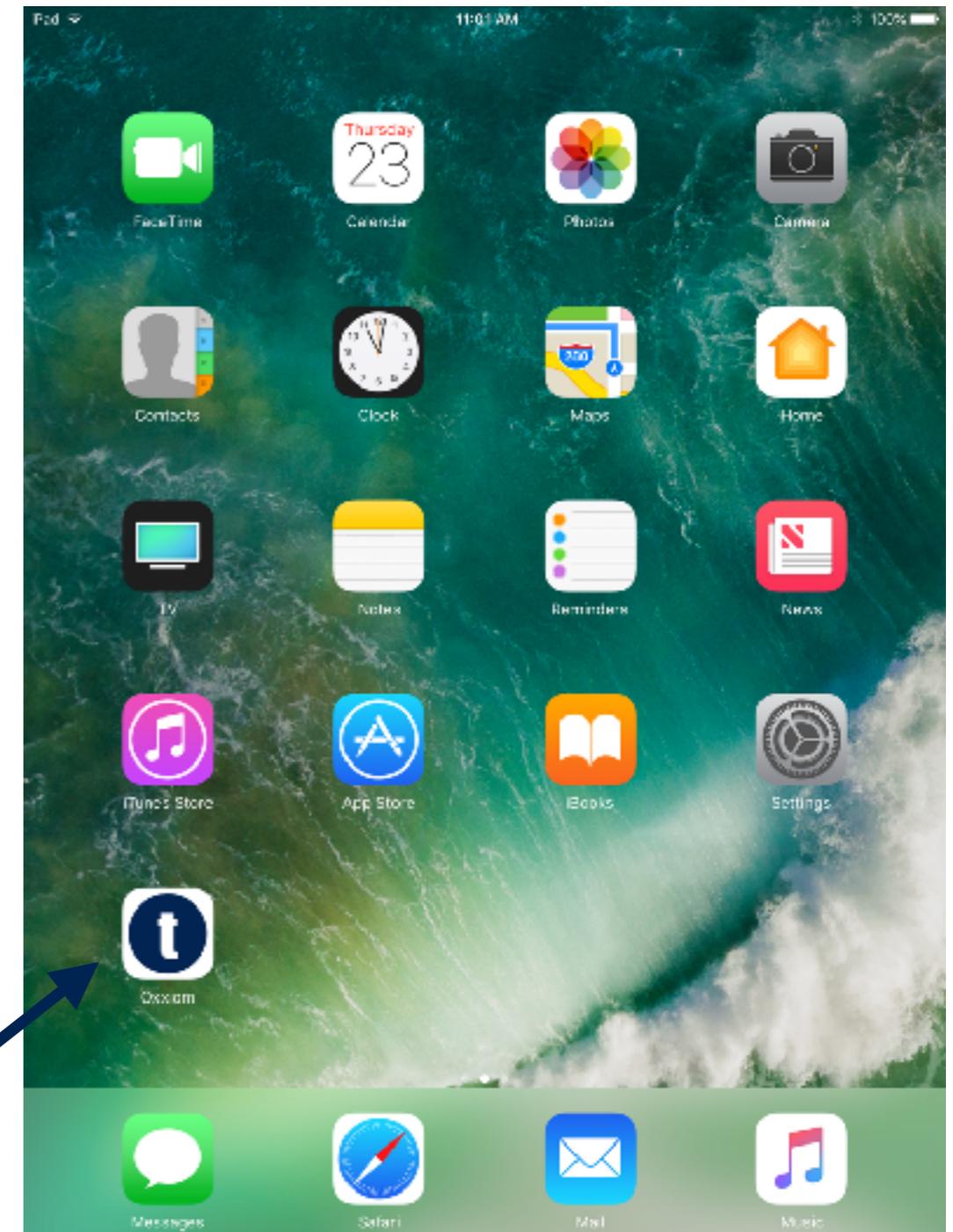
- Keep OXXIOM away from children and pets. OXXIOM is not a toy
- OXXIOM complies with part 15 of the FCC Rules. Operation is subject to the following two conditions: (1) OXXIOM may not cause harmful interference, and (2) OXXIOM must accept any interference received, including interference that may cause undesired operation. Any changes or modifications to OXXIOM not expressly approved by True Wearables may cause harmful radio frequency interference and void your authority to operate OXXIOM
- OXXIOM is a wireless device. Other equipments may interfere with OXXIOM even if they comply with CISPR emission requirements
- OXXIOM uses a non-rechargeable LiMnO<sub>2</sub> battery. Do not recharge, heat, expose to open flames, or disassemble OXXIOM
- OXXIOM must be used in conjunction with other methods of assessing clinical signs and symptoms (adjunctive device)
- Inspect OXXIOM measurement site at least every 8 hours to ensure correct sensor alignment and placement
- Patient sensitivity to OXXIOM biocompatible adhesive may vary due to skin condition
- Avoid excessive pressure to OXXIOM measurement site as this may cause damage to the skin beneath OXXIOM or inaccurate measurements
- No modifications to OXXIOM are allowed as they may affect OXXIOM performance
- Do not use a damaged OXXIOM. If OXXIOM is damaged, discontinue use immediately and replace OXXIOM unit
- OXXIOM general operation may be affected when in proximity to Electrosurgical Unit (ESU)
- Do not use OXXIOM in an Magnetic Resonance(MR) environment
- Do not use OXXIOM in an explosive atmosphere
- OXXIOM is not defibrillation proof (IEC 60601-1)
- OXXIOM has been tested and found to comply with IEC 60601-1-2 for electromagnetic compatibility of medical devices. This standard is designed to provide reasonable protection against harmful interferences in a typical medical installation. However, given the increased number of radio-frequency transmitting equipments and other interference sources in medical installations and other environments, high interference levels might occur and disrupt the operation of OXXIOM
- Follow governing ordinances and recycling instructions regarding disposal or recycling of OXXIOM and its components, including batteries
- OXXIOM contains WEEE materials. In compliance with the European directive on Waste Electrical and Electronic Equipment (WEEE) 2002/96/EC, do not dispose OXXIOM as unsorted waste. Please contact True Wearables regarding disposal or recycling of OXXIOM and its components.

# STEP 1: Download OXXIOM App

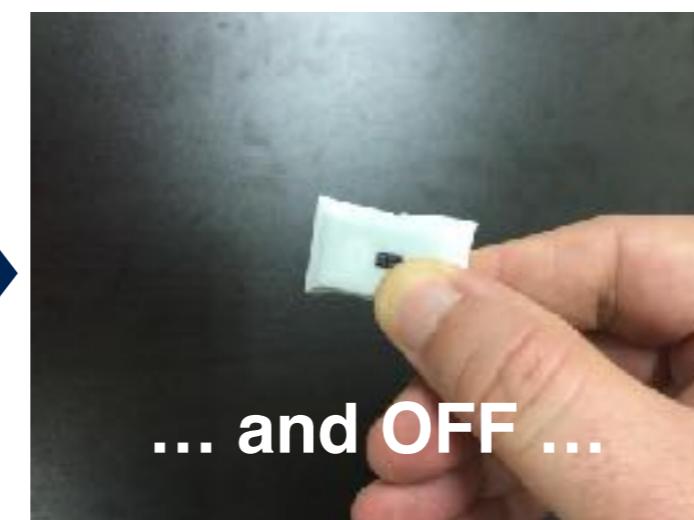
In your iPad or iPhone,  
select App Store  
and search for Oxxiom



Oxxiom App will be  
now available in your  
iPad or iPhone



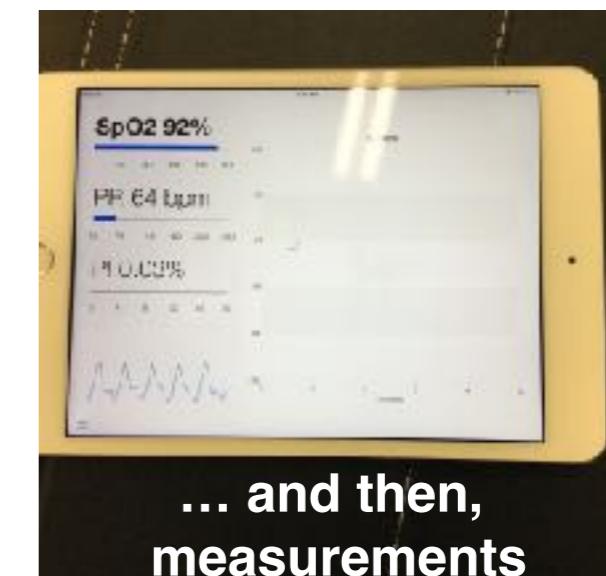
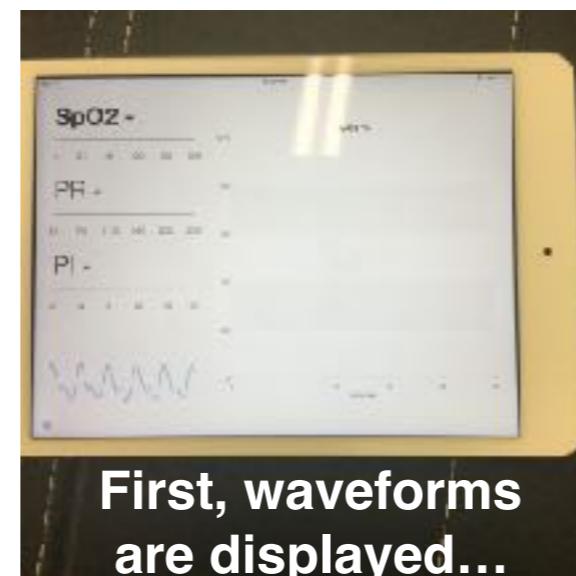
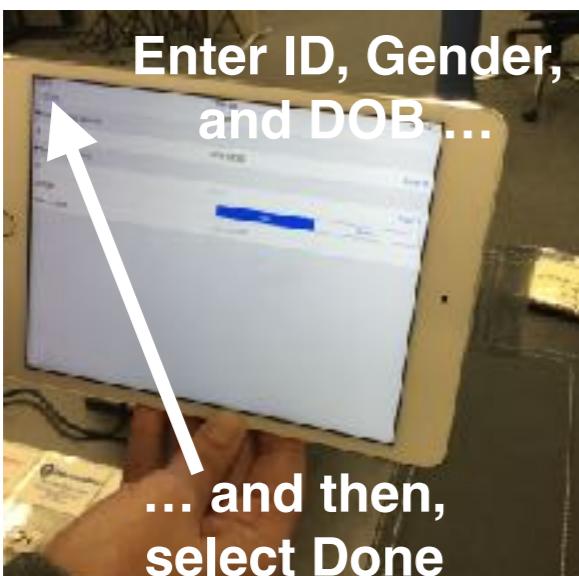
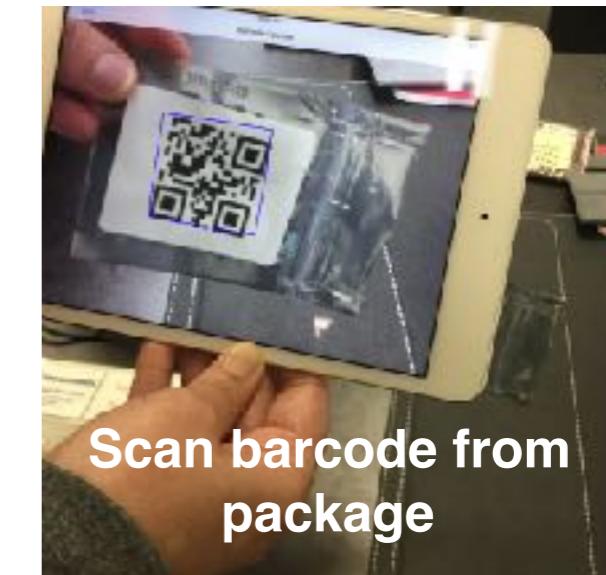
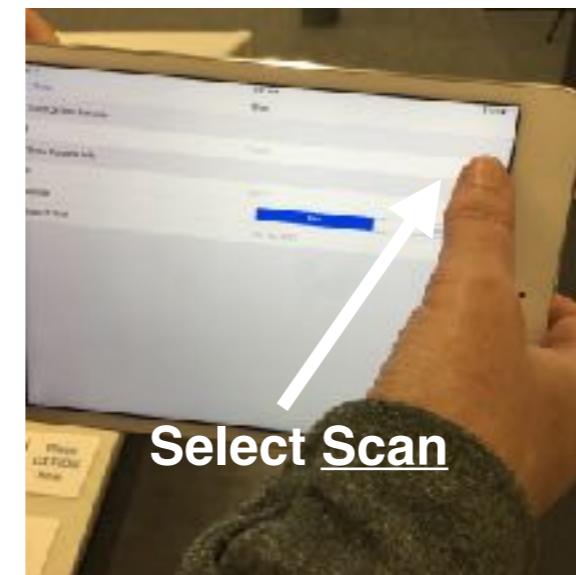
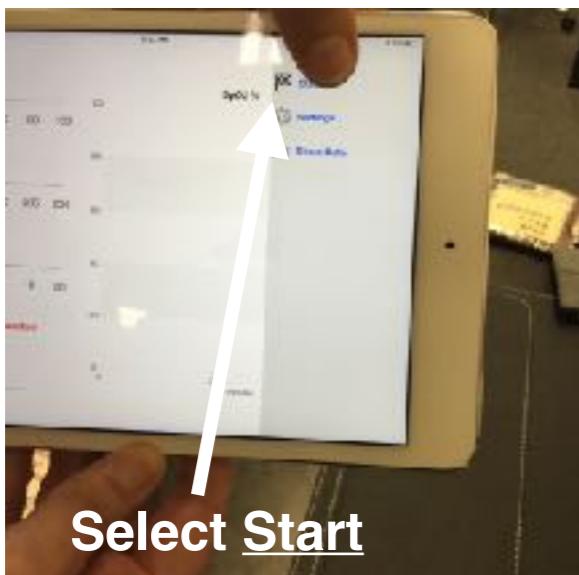
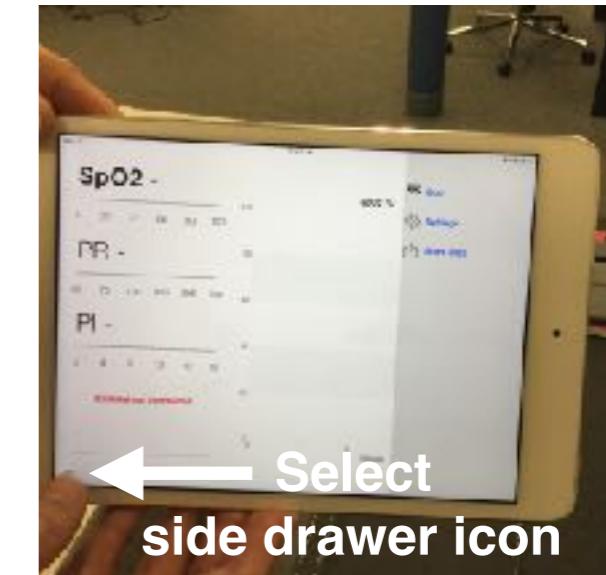
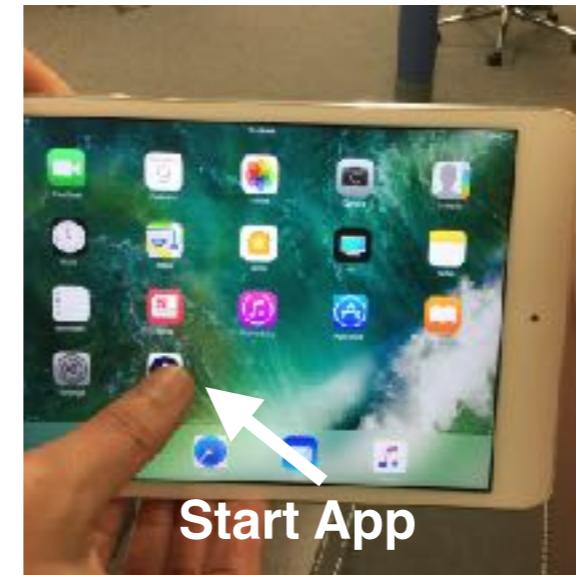
## STEP 2: Unpack and Turn On OXXIOM



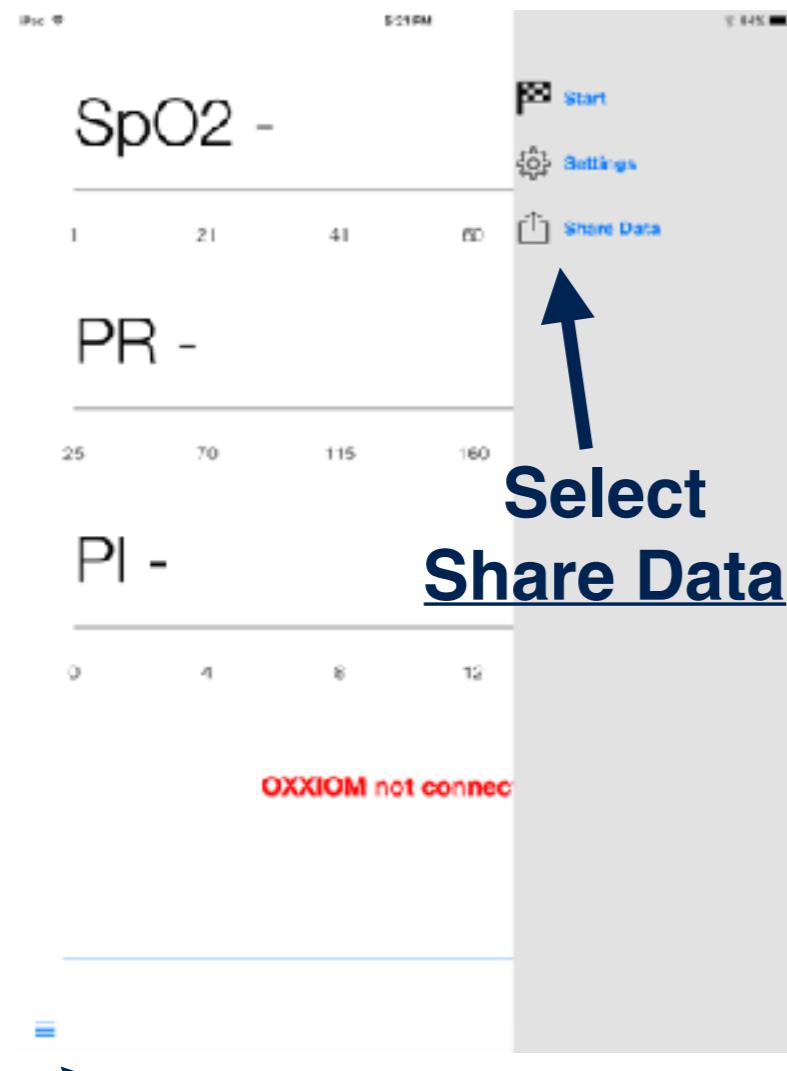
## STEP 3: Attach OXXIOM to Patient



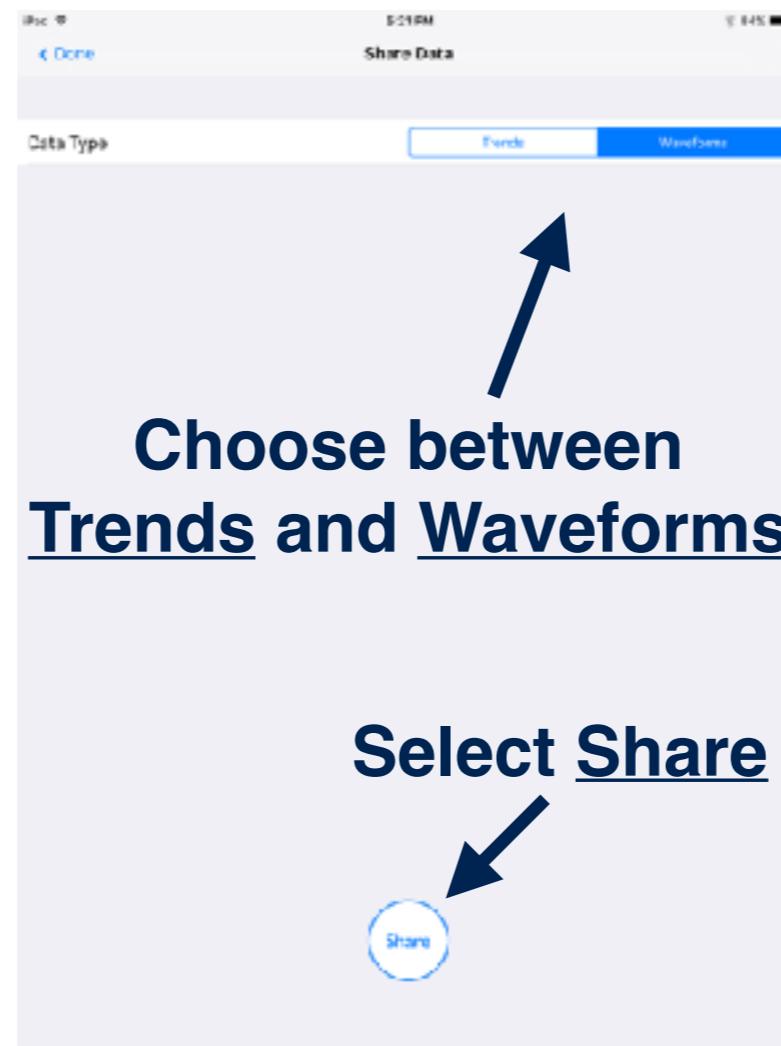
# STEP 4: Pair OXXIOM With App



# SHARING DATA

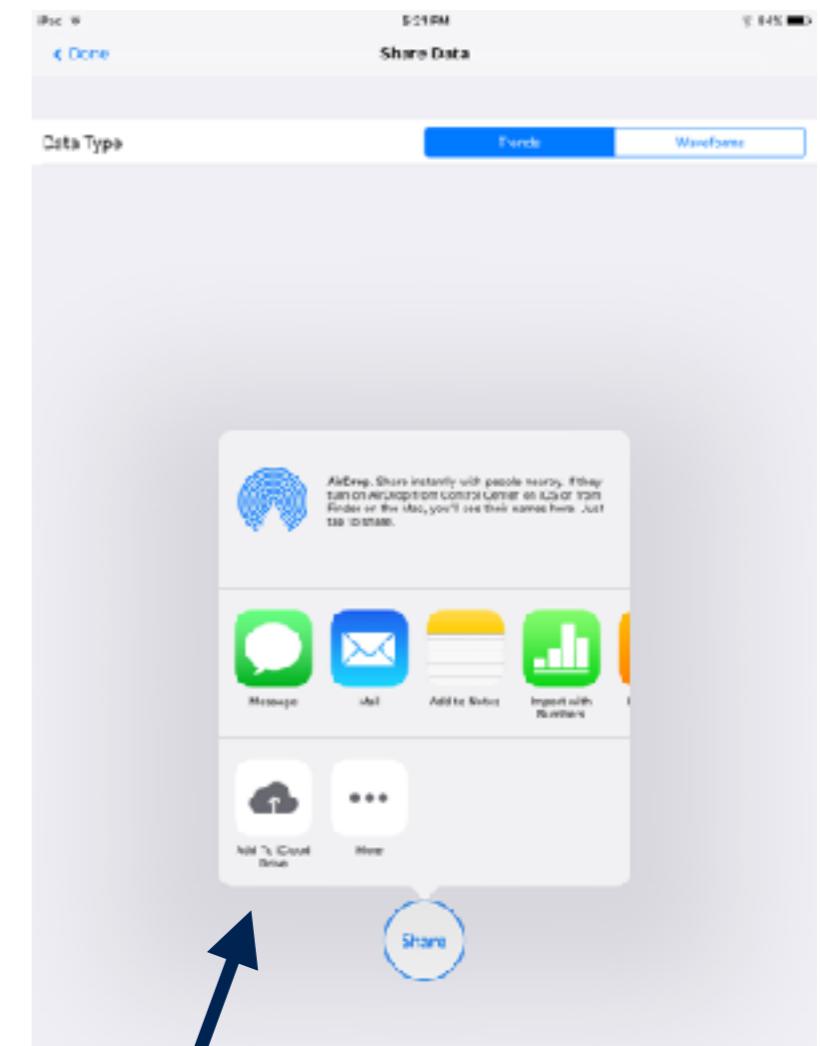


Select side drawer icon

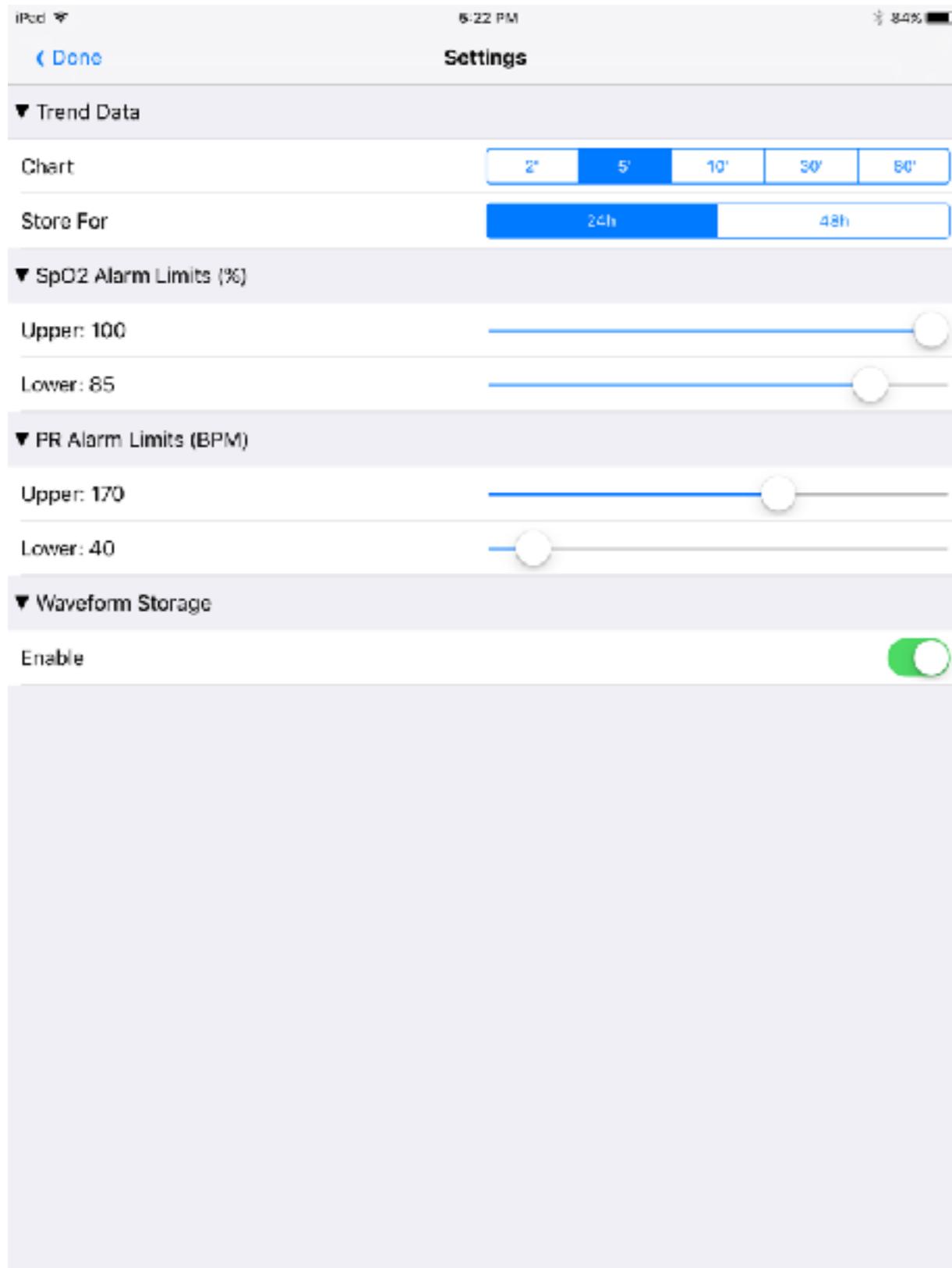


Select Share

Choose sharing method  
(Mail, Cloud, Message, AirDrop, etc.)



# SETTINGS



**Set memory trend data chart (landscape mode)**

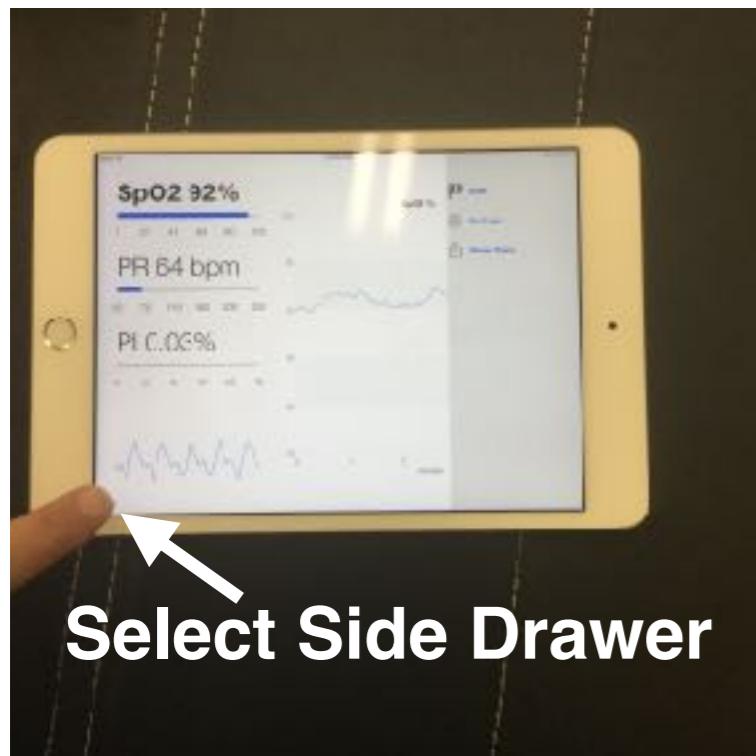
**Set storage time for trend data (Measurements)**

**Set upper and lower SpO2 alarm limits**

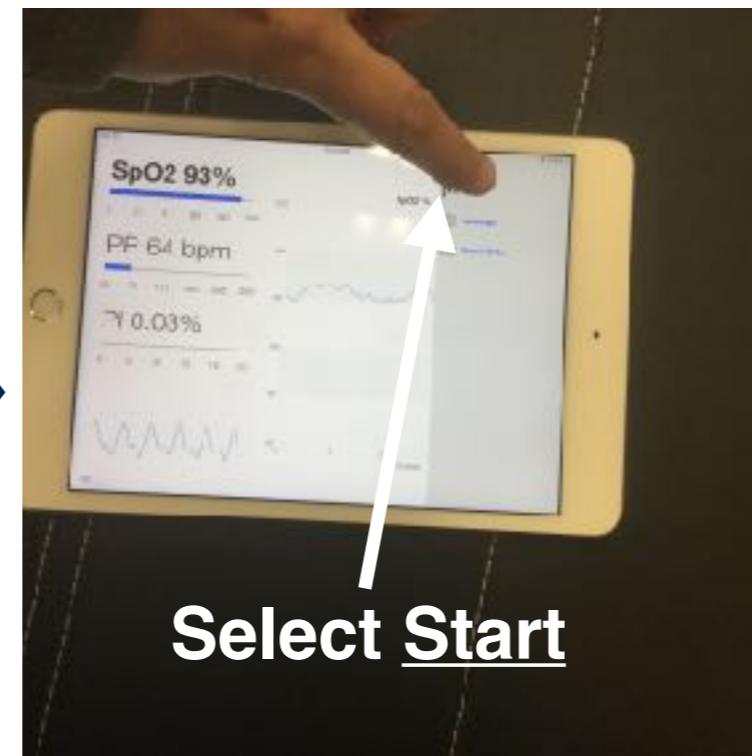
**Set upper and lower PR alarm limits**

**Set waveform storage ON or OFF**

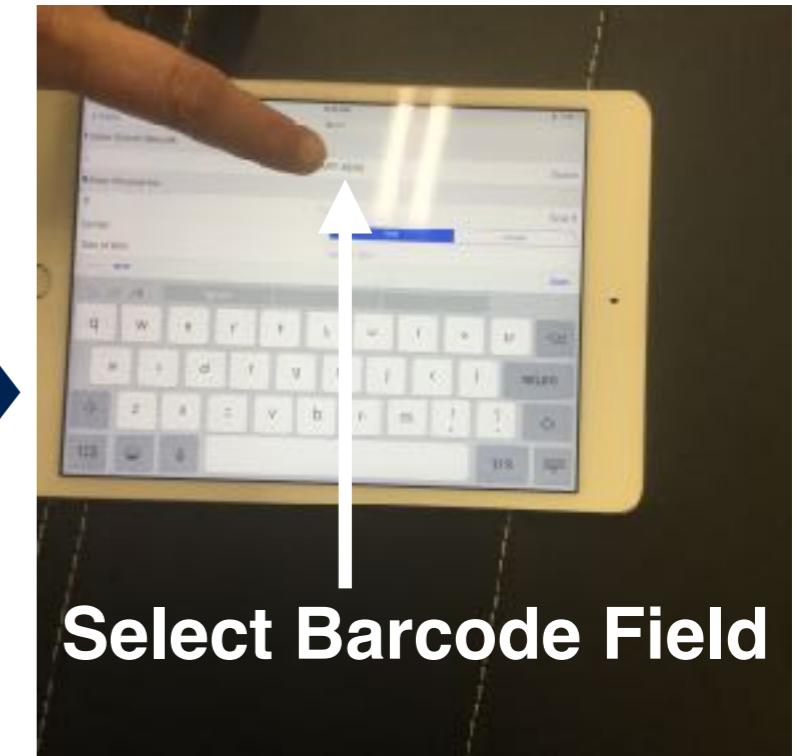
# UN-PAIRING OXXIOM



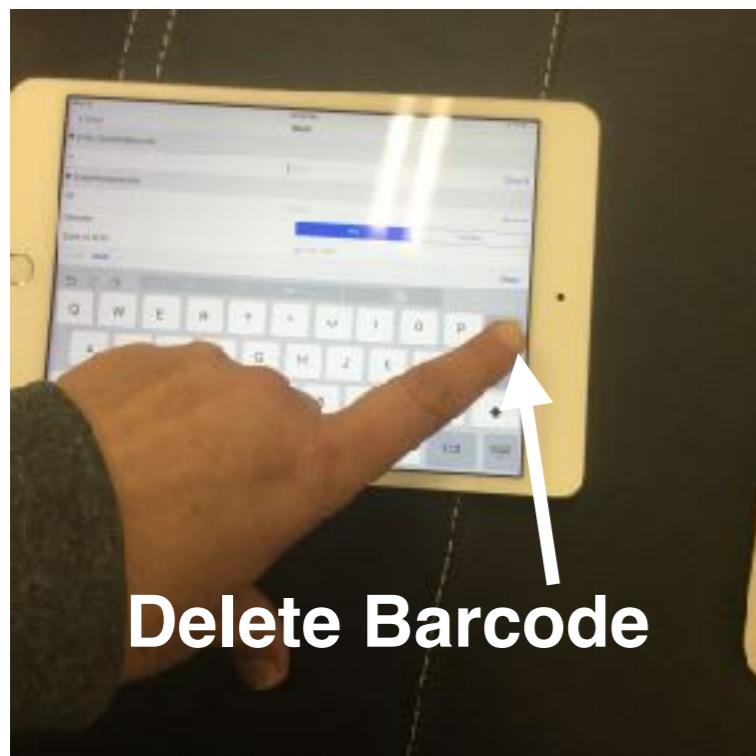
Select Side Drawer



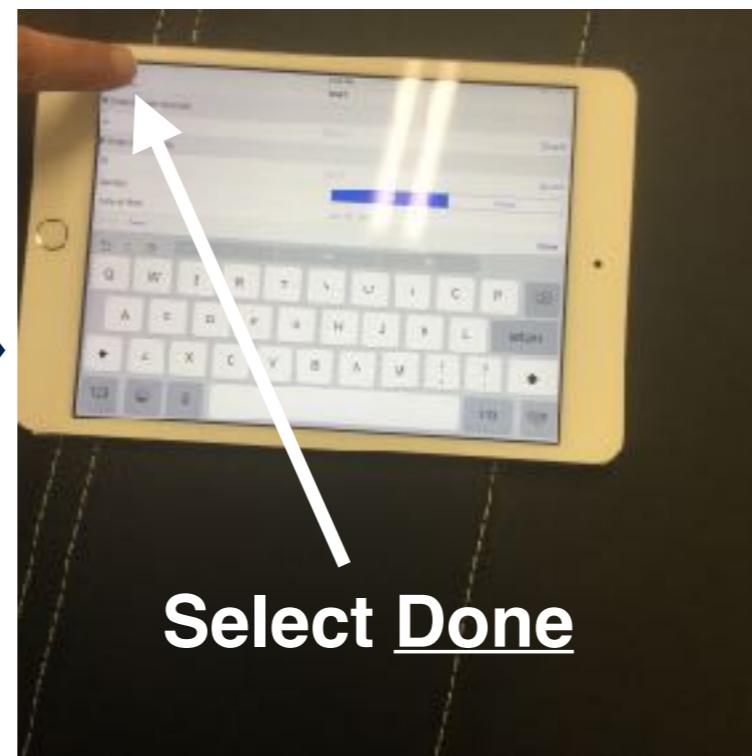
Select Start



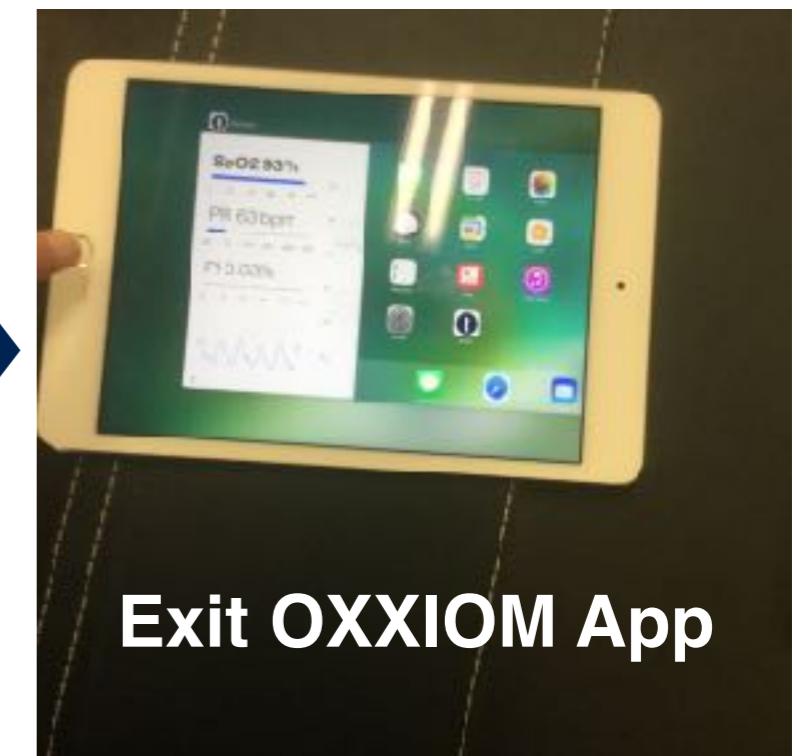
Select Barcode Field



Delete Barcode



Select Done



Exit OXXIOM App

# COPYRIGHT NOTICE

- Copyright © 2017 by True Wearables, Inc.
- This User Manual is intended to be employed only for the purpose of OXXIOM operation and reference. All rights reserved. No part of this Manual may be reproduced, distributed, or transmitted in any form or by any means, including photocopying, recording, or other electronic or mechanical methods, without the prior written permission of True Wearables, Inc.
- True Wearables intends to keep this Manual confidential and owns all rights to this unpublished work. True Wearables reserves the right to make changes or to discontinue this Manual and OXXIOM without notice or obligation

# CONTACT INFORMATION

*At True Wearables, we value our customers and take great pride in our products and technology. We welcome your inquiries and comments. Please direct all inquiries and/or comments regarding OXXIOM by email to:*

**contact@truewearables.com**

*or by mail to:*

**True Wearables, Inc  
29826 Avenida de Las Banderas, Suite 300  
Rancho Santa Margarita CA 92688**

*with message subject:*

**Customer Service, OXXIOM**

*and message with inquire and/or comments*

# OXXIOM LABEL INFORMATION



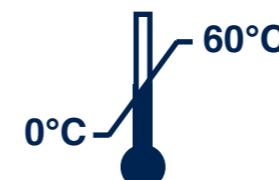
OXXIOM should not be used on a sterile field and should not be sterilized



Not made with natural rubber latex



Consult User Manual for instructions on how to use OXXIOM



Storage temperature range: 0°C to 60°C

IPX1

Resistant to liquid ingress



Do not re-use



Type B Applied Part  
(Comply with specified requirements for protection against electric shock)



Mark of Conformity to European Medical Device Directive 93/42/EEC

# FCC DECLARATION

This equipment has been tested and found to comply with the limits for a Class B 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:

- Reorient or relocate the receiving antenna
- Increase the separation between the equipment and receiver
- Connect the equipment into an outlet on a circuit different from that to which the receiver is connected
- Consult the dealer or an experienced radio/TV technician for help