TROUBLESHOOTING

			TROBELLY WITH THIS BESTOCK
An alarm is sounding, indicating an abnormal oxygen saturation or heart rate.	Your finger is not inserted correctly into the device.	Insert your finger until the tip of your finger touches the end of the device and try measuring again.	EXPLOSION HAZARD — DO NOT USE THE FINGERTIP PULSE OXI- METER IN A FLAMMABLE ATMOSPHERE WHERE CONCENTRATIONS OF FLAMMABLE ANESTHETICS OR OTHER MATERIALS MAY OCCUR.
	Your oxygen saturation is abnormal (under 94%).	Try measuring again a couple of times. If the alarm keeps beeping, get in contact with	DO NOT THROW BATTERIES IN A FIRE AS THIS MAY CAUSE THEM TO EXPLODE.
		your doctor for further advice.	DO NOT ATTEMPT TO RECHARGE NORMAL DRY-CELL
	Your heart rate is abnormal (under 50 or above 130).	Try measuring again a couple of times. If the alarm keeps	BATTERIES, THEY MAY LEAK. AND MAY CAUSE A FIRE OR EVEN EXPLODE.
, ,		beeping, get in contact with your doctor for further advice.	THE BATTERIES MUST BE TAKEN OUT FROM THE BATTERY COM- PARTMENT IF THE DEVICE WILL NOT BE USED
The oximeter does not power	er pressing the button. a fresh pair of AAA batteries.	Replace the old batteries with	FOR A LONG TIME.
on after pressing the button.		DO NOT USE THIS DEVICE IF THE BATTERY COVER IS OPEN.	
incorrectly.	The batteries are installed incorrectly.	Make sure the batteries are installed correctly, with the plus-sides of both batteries facing down.	THE BATTERIES MUST BE PROPERLY DISPOSED OF ACCORDING TO LOCAL REGULATION AFTER THEIR USE.
	The device is broken.	Contact the iProven service center at iproven.com/pages/support for	KEEP THE OPERATING ENVIRONMENT FREE OF DUST, VIBRATIONS, CORROSIVE, OR FLAMMABLE MATERIALS, AND EX- TREMES OF TEMPERATURE AND HUMIDITY.
		help.	DO NOT OPERATE THE UNIT IF IT IS DAMP OR WET BECAUSE OF
The screen suddenly turns off.	The automatic shutoff feature turns off the device after 16 seconds of	No need to do anything.	CONDENSATION OR SPILLS. AVOID USING THE EQUIPMENT IM- MEDIATELY AFTER MOVING IT FROM A COLD ENVIRONMENT TO A WARM, HUMID LOCATION.
inactivity.		NEVER USE SHARP OR POINTED OBJECTS TO OPERATE THE	
	The batteries are depleted.	Replace the old batteries with	FRONT-PANEL SWITCHES.
		a fresh pair of AAA batteries.	THE DEVICE SHOULD BE KEPT AWAY FROM CHILDREN AND PETS TO AVOID SWALLOWING.

SPECIFICATIONS

DO NOT REPAIR OR MODIFY THIS DEVICE YOURSELF, PLEASE CONTACT IPROVEN CUSTOMER SUPPORT IF THERE IS A PROBLEM WITH THIS DEVICE.

XPLOSION HAZARD — DO NOT USE THE FINGERTIP PULSE OXI-METER IN A FLAMMABLE ATMOSPHERE WHERE CONCENTRATIONS OF FLAMMABLE ANESTHETICS OR Dimensions:74 mm (L) x 37mm (W) x 38mm (D) OTHER MATERIALS MAY OCCUR. Weight -approx: 70g±2g (including 2 x AAA battery)

DO NOT THROW BATTERIES IN A FIRE AS THIS MAY Classification CAUSE THEM TO EXPLODE.

Anti-electric Shock Type: Internally powered equipment DO NOT ATTEMPT TO RECHARGE NORMAL DRY-CE! BATTERIES, THEY MAY LEAK, AND MAY CAUSE A FIRE Anti-electric Shock Degree: Type BF equipment

Physical Characteristics

EMC: Type B THE BATTERIES MUST BE TAKEN OUT FROM THE BATTERY CO

PARTMENT IF THE DEVICE WILL NOT BE USED Mode of operation: Continuous Operation

Enclosure Degree of ingress protection: IP22

The Fingertip Pulse Oximeter Specifications:

DO NOT USE THIS DEVICE IF THE BATTERY COVER IS OPE IP22 means shell of this product can with

select dropping to the surface.

KEEP THE OPERATING ENVIRONMENT FREE OF DUST. IBRATIONS, CORROSIVE, OR FLAMMABLE MATERIALS, AND EX-TREMES OF TEMPERATURE AND HUMIDITY.

O NOT OPERATE THE UNIT IF IT IS DAMP OR WET BECAUSE OF ONDENSATION OR SPILLS, AVOID USING THE EQUIPMENT I Power Consumption MEDIATELY AFTER MOVING IT FROM A COLD ENVIRONMENT TO

NEVER USE SHARP OR POINTED OBJECTS TO OPERATE THE FRONT-PANEL SWITCHES. Environmental:

THE DEVICE SHOULD BE KEPT AWAY FROM CHILDREN AND PETS TO AVOID SWALLOWING

SPECIFICATIONS

mperature:	−25°C to 55°C
umidity:	15% to 93% non-condensing
re	70Kpa-106Kpa

Electronics Parameters:

Relative H

Hemoglobin saturation

Pulse rate:

thstand the water from any		Hemoglobin display	saturation	35-100%	
		Pulse rate Di	splay	30-250 BPM	
lkaline battery		Resolution	Hemoglobin Saturation	1%	
		l			

2xAAA 1.5v al

erature:	5°C to 40°C		
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SPECIFICATIONS

LABELING OR PACKAGING

Accuracy

Probe LED Specification:

−25°C to 55°C		Display
15% to 93% non-condensing	PI	Resolution
70Kpa-106Kpa	FI	Measure

Sound Reminder Limit default value:

,			Wave Lengt
limit: 100/ bottom limit:94		RED	660±2 nm
11 1: 120 1 11 1: 50	1	RED	000±2 mm
limit: 130 /bottom limit:50		L.C. DED	005 : 2

	Value	
ation	35-100%	Symbol
	30-250 BPM	★
noglobin ıration	1%	LOT
se rate	1 BPM	
noglobin aration	±2% (90%-100%)	
mation	±3% (70%-90%), Unspecified (<70%)	SN
se rate	+1 RPM	

6	_	***	Information of manufacture, includin name and address
		1	Temperature limitation
: 0.1% 6: 1%		\A	When the end-user wishes to discard thi product, it must be sent to separat

				_	recycling
	Wave Length	Radiant Power			recycling
	wave Length	Radiant I ower			T. 11
	660±2 nm	1.8 mW			Follow instruction for use
	005:3	100 111			
)	905±2 nm	$2.0 \mathrm{mW}$		IP22	Anti-dust & Anti-water class
SYMBOLS ON THE MONITOR,			IFZZ	Anti-dust & Anti-water class	
		1	*Batch code Date of m	anufacturer and Serial No are pri	

*Batch code. Date of manufacturer and Serial No are printed on the label on the battery cover

Symbol	Description	Manufacturer's Declaration of the EMC Guidance and manufacturer's declaration				
*	Type BF Equipment		1	Guidance an electromagnetic	d manufa	SYSTEMS acturer's
LOT	Batch code*	2	,	The A330 Puls electromagnetic or the user of A	e Oximeter environment	specified be
П		1		used in such an er	nvironment.	
\sim	Date of manufacture*		3	Emissions test	Complian-	Electroma
		l Ľ	1	Limssions test	ce	environme
SN		Ι Γ	T	RF emissions		The
SN	Serial NO*	4	4		Group 1	Oximeterus
			- 1	CISPR 11		for ite

n of manufacture, including	Γ			Therefore, its RF emissions	Immunity	IEC 60601
address				are very low and are not likely to cause any	test	test level
re limitation		RF emissions		interference in nearby electronic equipment.	Electrostatic	
end-user wishes to discard this it must be sent to separate		CISPR 11	Class B	The A330 Pulse	discharge (ESD)	± 8 kV contact
facilities for recovery and		Harmonic emissions	N/A	for use in all establishments, including domestic establish	IEC 61000-4-2	± 15 kV air
	L	IEC 61000-3-2		ments and those directly		
struction for use		Voltage fluctuations /		connected to the public low-v oltage power supplynetwork t hat supplies buildings used fo		± 2 kV for
& Anti-water class	1	7 flicker	N/A	r domestic purposes.	Electrostatic	nower loi

collection facilities for recovery

manufacturer's declaration - electromagnetic

nufacturer's	declarati	on –	immunity -for all EQUIPMENT and SYSTEMS
ent specified	ded for use below. The d hould assure t	customer	immunity The A330 Pulse Opiniotes is intended for use in
	magnetic		in such an environment.
The	ment - guida A330	Pulse	
THE	11000	i disc	

Therefore, its RF emissions	Immunity	IEC 60601	Con
are very low and are not ikely to cause any nterference in nearby	test	test level	nce
Plectronic equipment. The A330 Pulse Dximeter is suitable	Electrostatic discharge (ESD)	± 8 kV contact	± 8
for use in all establishments, neluding domestic establish nents and those directly	IEC 61000-4-2	± 15 kV air	± 1: air
connected to the public low-v oltage power supplynetwork t nat supplies buildings used fo domestic purposes.	Electrostatic transient /	± 2 kV for power supply lines	

61000-4-4 Guidance and manufacturer's declaration - electromagnetic

mulacturer's declaration =	immunity -for all EQUIPMENT and SYSTEMS	
eter is intended for use in the ent specified below. The customer	Guidance and manufacturer's declaration – electromagnetic immunity	
e Oximeter should assure that it is	The A330 Pulse Oximeter is intended for use in telectromagnetic environment specified below. The customer	
an- Electromagnetic	the user of the A330 Pulse Oximetershould assure that it is used in such an environment.	

Immunity	IEC 60601	Complia	Electromagnetic
test	test level	nce level	environment - guidance
Electrostatic discharge (ESD) IEC 61000-4-2	± 8 kV contact ± 15 kV air	± 8 kV contact ± 15 kV air	are covered with synthetic material,
Electrostatic transient / burst	± 2 kV for power supply lines	N/A	Mains power qu should be that of a typical commercia

test level nce level environment s should be wood, amic tile. If floors If the user of the thetic material, supply uld be at least mains interruptions, 0. the A330 Pulse from an uninterruptible power supply ıld be that of a

> magnetic characteristic of typical location in typical commercial NOTE 1. At 80 MHz and 800 MHz, the higher frequency range applies. hospital environmen NOTE 2. These guidelines may not apply in all situations. Electromagnetic is affected by absorption and reflection from structures, objects and people. NOTE UT is the a. c. mains voltage prior to application of the

magnetic fields should

between

environment electromagnetic immunity -for EQUIPMENT Mains power quality SYSTEM that are not LIFE-SUPPORTING should be that of a

Immunity The A330 Pulse Oximeter is intended for use in the electromagnetic environment specified below. The customer or the user of the A330 Pulse Oximeter should assure that it is used in such an environment.			d below. The customer or should assure that it is used		transmitters, an electromagnetic site survey should be considered. If the measured field strength in the location in which the A330 Pulse Oximeter is used exceeds the applicabl RF compliance level above, the A330 Pulse Oximeter should be
Immunit			Electromagnetic		observed to verify normal operation. If abnormal performance is
y test	test level	nce level	environment - guidance		observed, additional measures may be necessary, such a
Conduct	3 Vrms	N/A	Portable and mobile RF		
ed RF			communications	the transmitter in watts	reorienting or relocating the A330 Pulse Oximeter.
	150 kHz to 80		equipment should be	(W) according to the	b Over the frequency range 150 kHz to 80 MHz, field
IEC	MHz		used no closer to any part	transmitter manufacturer	strengths should be less than 3V/m.
61000-4-			of the A330 Pulse	and d is the	suchgue should be less than 5 V/III.
6	6Vrms in	10 V/m	Oximeter, including	recommended separation distance in metres (m).b	Recommended separation distances between portable

the equation applicable to determined by

E1 NP 80 MHz to 800 marked with

than the compliance le

in each frequency range.

Interference may occur

the frequency of the electromagnetic

transmitter.

Recommended

separation distance

where p is the maximum

A330 Pulse Oximeter

electromagnetic environment in which radiated RF disturbar

are controlled. The customer or the user of the A330 Pul Oximeter can help prevent electromagnetic interference by maintaining a minimum distance between portable and mobile RF communications equipment (transmitters) and the A330 Pulse Oximeter as recommended below, according to the maximum output power of the communications equipment

Separation distance according to frequency of

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 T broadcast cannot be predicted theoretically with accuracy. assess the electromagnetic environment due to fixed R

For transmitters rated at a maximum output power not listed above the recommended separation distance d in metres (m) can

and mobile. RF communications equipment and the EQUIPMENT or SYSTEM -for EQUIPMENT and SYSTEMS that are not LIFE-SUPPORTING.

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.



be estimated using the equation applicable to the frequency of

the transmitter, where P is the maximum output power rating o

the transmitter in watts (W) according to the transmitter

NOTE 1 At 80 MHz and 800 MHz, the separation distance fo

WAREHOUSE Beaverton, Oregon 9450 SW Gemini D Beaverton, OR 97008-7105



HOW TO USE THE OXI-33



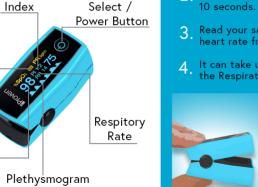
WHAT'S IN THE BOX?

- Oximeter iProven OXI-33
- Two AAA-batteries
- Lanyard
- Oximeter Pouch
- Instruction Manual



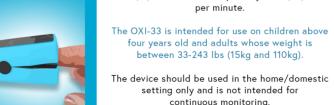
COMPONENTS OF THE DEVICE

Perfusion



Intended Use HOW TO TAKE A MEASUREMENT

- Open the clip and place your finger for non-invasive measuring and monitoring of the inside the oximeter. Make sure your oxygen saturation and pulse rate. The digital nail touches the upper part of the cli
- blood by checking the hemoglobin levels in your Press the button and wait for
- Read your saturation level and heart rate from the display. beats per minute, and displays a bar graph that
- It can take up to 1 or 2 minutes before the Respiratory Rate is displayed.



MEASUREMENT PRINCIPLE

The iProven Fingertip Pulse Oximeter is intended The fingertip pulse oximeter contains a icroprocessor and a display. The displa shows the oxygen saturation, the pulse device measures the oxygen saturation of your rate and the intensity of your heart beat.

arterial blood. The monitor detects signs of low levels of oxygen in the body. It measures the pulse rate in TT-1-3

illustrates the strength of the pulse signal.

The monitor also displays the Perfusion Index (PI) in % and the Respiratory Rate (RR) The display continuously gives information about the level of oxygen in you<u>r body and</u>

> your pulse rate per minute. e monitor updates its calculations requ ring use to show immediate readings. '

> > monitor starts to beep if the value of the

oxygen saturation 94% or lower. It will also beep when your heartbeat is below 50 or above 130.

PLEASE NOTE

into the device

on the same arm

or is in shock

applications

Measurements may be inaccurate in the following situations: When you're moving your fingers In a very cold environment

- When your finger isn't placed properly (hb) attaches oxygen to the red blood cell and carries it through the body. 1 hemoglobin mole
- When using a Blood Pressure Monitor/Cuff
- In case someone has hypotension, severe vasoconstriction, severe anemia or hypothermia
- In case someone has a cardiovascular arrest
- In case you have enamel or acrylic fingernail polish, false fingernails or other fingernail
- Excessive ambient light, such as a fluorescent
- the blood flow in the extremity or body part lamp or direct sunlight, may affect the result. where it is measured. A PI is considered hea
- prolonged monitoring, do not forget to check blood flow and skin condition regularly, at least every 2 hours. If necessary, change the measuring site to a finger of the other hand.
- Do not use the Fingertip Pulse Oximeter in an MRI or CT environment you measure, the more accurate the RR become
- For sport and aviation use only

Q: HOW DO I CLEAN THE DEVICE?

- WHEN POWERING ON THE MONITOR. THE LED DISPLAY

A: PUT THE THIN END OF THE ROPE THROUGH THE HOLE

Q WHY DOES MY RESPIRATORY RATE (RR) NOT DISPLAY

THEN PUT THE THICK END OF THE POPE THROUGH THE LOOP

A IT MAY TAKE UP TO 1 OR 2 MINUTES BEFORE YOUR RR CAN BE

A THIS DEVICE IS A CONTINUOUS MONITORING DEVICE. WHEN

DISPLAYED. THE DEVICE NEEDS 1 OR 2 MINUTES TO GIVE AN

Q WHY DO THE VALUES OF MY OXYGEN SATURATION, RESPI

ONE OF THE MEASURED VALUES CHANGES. THIS WILL BE

Oxygen saturation indicates how much oxyge the blood is carrying as a percentage of the

A: YOU CAN USE MEDICAL ALCOHOL TO DISINFECT I maximum it could carry. Red blood cells conta

PLEASE LISE A SOFT CLOTH TO CLEAN THE WHOLE LINIT ron-containing protein hemoglobin. Hemoglo DON'T USE ANY ABRASIVE OR VOLATILE CLEANERS Q: HOW CAN I REPLACE THE BATTERIES?

DOESN'T LIGHT UP.

ACCURATE INDICATION.

DISPLAYED ON THE SCREEN.

Q: HOW DO I ATTACH THE LANYARD?

AND PULL THE END TO SECURE IT.

A: OPEN THE BATTERY COVER. cule can attach up to 4 oxygen molecules. If a INSERT THE BATTERIES ACCORDING TO THE POLARITY the hemoglobin molecules are carrying 4 oxyg INDICATIONS. CLOSE THE BATTERY COVER.

Q: WHEN DO I REPLACE THE BATTERIES molecules, it is said that you have a saturat

- A: LOW BATTERY SIGN SHOWS ON THE DISPLAY. of 100%. A healthy person with healthy lung THE LED DISPLAY DIMS.
- breathing air at sea level, will have an oxyge
- saturation of between 95% and 100%.

WHAT IS PERFUSION INDEX?

Perfusion Index or PI indicates the strength of

WHAT IS OXYGEN SATURATION?

· When someone needs this oximeter for between 0.3-20%. WHAT IS RESPIRATORY RATE?

> Respiratory Rate or RR reflects the number of breaths a person takes per minute. The longer

CHANGE PARAMETER SETTINGS

1.) REMINDER SETUP

2.) LIMIT SETUP

MENU. MAKE SURE THE * IS ON TOP OF THE MENU AN

PRESS THE 'ON' BUTTON A LITTLE LONGER.

Remind Setup Sound Reminder

Brightness

BY ENTERING THE MENU OF THIS OXIMETER, YOU CAN CHANGE SOME OF THE PARAMETER SETTINGS. TO ENT INDICATED LEVEL. HE PARAMETER SETTINGS, SWITCH ON THE DEVICE A BEEP: SWITCH THE BEEP ON OR OFF WHEN YOU

PRESS THE 'ON' BUTTON AGAIN. THERE ARE 2 SUBMENU'S:

> SELECT MODE: SELECT IF YOU CLIP THI OXIMETER ON THE FINGER OF YOUR RIGHT (R) OR LEFT (L) HAND, OR IF YOU WANT TO USE IT

RESTORE: SELECT THIS IF YOU WANT TO RESET THE

BRIGHTNESS: ADAPT THE BRIGHTNESS OF THE DISPLAY, VALUES RANGE FROM 1-5.

LIMIT MENU

/-: INCREASES (+) OR REDUCES (-) THE NUMBERS

REMINDER MENU SOUND REMINDER: STARTS TO BEEP IF YOUR SPO

LEVELS OR HEART RATE ARE ABOVE THE HIGHES INDICATED LEVEL OR BELOW THE LOWEST

START UP THE DEVICE.

VERTICALLY (V).

FACTORY SETTINGS.

YOU CAN SCROLL THROUGH THE MENU BY PRESSING SET THE UPPER AND LOWER LEVEL FOR YOUR SPO2 AND THE 'ON' BUTTON, THE * WILL INDICATE WHERE YOU PULSE RATE, IF YOU SET 'SOUND REMINDER' TO 'ON' IN ARE IN THE MENU, WHEN YOU WANT TO CHANGE THE REMINDER MENU. THE OXIMETER WILL START TO SETTING, YOU CAN PRESS THE 'ON' BUTTON A LIT BEEP IF IT MEASURES VALUES THAT ARE LOWER OF LONGER, AND THE VALUE WILL CHANGE, WHEN YOU HIGHER THAN THE VALUES THAT ARE SET. VANT TO SWITCH BETWEEN THE 'REMINDER' AND 'LII

WHEN SETTING THE SPO2 OR PULSE RATE.

TROUBLESHOOTING

Insert your finger until the tip of

your finger touches the end of th

device and try measuring again

Do not run or walk around wi

measuring. Hold your hands st when taking a measurement.

screen.		
Perfusion Index does not show up on screen.	Your finger is not inserted correctly into the device.	Insert your finger until the tip of your finger touches the end of the device and try measuring again.
	Your perfusion (strength of your blood flow) is too low to be measured.	Try measuring a couple more times. Insert your finger until your fingeript buches the end of the device. If it keeps happening and you feel other symptoms of poor blood circulation like numbness, tingling or pain, contact your doctor for advice.
Oxygen saturation or heart rate is unstable.	Your finger is not inserted correctly into the device.	Insert your finger until the tip of your finger touches the end of the device and try measuring again. Make sure the device is shut snugly against the top and bottom of your finger. Do not curl your finger in the device.
	Your finger is trembling or shaking.	Put your hand with the device on a flat surface, like a table, to avoid trembling hands and fingers.