



EnsoSleep PPG Device Profile

Reference Document

Introduction

EnsoSleep PPG processes physiological data obtained during sleep studies and applies a variety of analytical approaches to identify the occurrence of sleep events that relate to the presentation of sleep disorders. At the end of the analysis, the system generates a score report that includes tables and graphs typical of those generated following manual scoring of sleep studies by certified technologists. The results of the automated scoring may be displayed using commercially available sleep study viewer applications or it may be displayed within this device; both allow for manual editing of the results and generation of a revised score report.

It is a stand-alone software device, which can be accessed from a computer with a compatible operating system platform and operated without being attached to any other physical device or any direct or indirect contact with a patient.



Please read this document carefully before using the device. User information can be found in the instructions for use available within the device.

Indications for Use

EnsoSleep PPG is a Software as a Medical Device (SaMD) that establishes sleep quality. EnsoSleep PPG automatically analyzes, displays, and summarizes Photoplethysmogram (PPG) data collected during sleep using compatible devices. EnsoSleep PPG is intended for use by and by order of a healthcare professional to aid in the diagnosis of sleep disorders including sleep apnea in adults.

The EnsoSleep PPG output, including automatically detected respiratory events and parameters, may be displayed and edited by a qualified healthcare professional. The EnsoSleep PPG output is not intended to be interpreted or clinical action taken without consultation of a qualified healthcare professional.

EnsoSleep PPG is not intended for use with polysomnography devices.

Contraindications

There are no known contraindications.

Target Population

EnsoSleep PPG is indicated for use in adults 22 years old and above.

Product Requirements

Computer Requirements:

- Operating System – Microsoft Windows Operating System version 10+
- Minimum 4 GB RAM, 16 GB RAM recommended
- Web Browser – Google Chrome (Mac or Windows) or Microsoft Edge (Windows)
- High Speed Internet connection – 12 Mbps or more recommended
- Patient Data File Formats – EDF or EDF+, HL7

EnsoSleep PPG is compatible with FDA-cleared medical purpose pulse oximetry devices that meet the data acquisition characteristics in this table:

Minimum Device Specification	Photoplethysmography	SPO2	Pulse /Heart Rate
Sampling Rate	16 Hz or greater	1 Hz	1 Hz
Digital Resolution	8 bit	n/a	n/a
Measurement Range	n/a	0 - 100%	30-250 BPM
Accuracy Range (70-100%)	n/a	+/- 3%	+/- 2%

Event Type	Event Subtypes	AASM Required Signals - minimum sampling rate; anatomical derivation, default settings
Respiratory Events	Sleep Disordered Breathing (Apneas or Hypopneas)	Photoplethysmogram (16 Hz) and Oximetry (1 Hz; Finger)
Sleep-Wake Events	Wake, Light Sleep, Deep Sleep, REM	Photoplethysmogram (16 Hz)

For a good user experience, EnsoData has identified the following as sources that have data acquisition characteristics and file formats meeting the system requirements:

- EDF files containing PPG data collected during sleep that are consistent with the acquisition characteristics mentioned above.
- Viatom Checkme O2

The above list is provided for informational purposes only and is correct at the time of issue; the list will be updated as additional data sources are identified and allowed by EnsoData. If your data source is not listed or does not meet the above acquisition characteristics, please email: support@ensodata.com.

Study Scoring

The autoscoring algorithm is based directly on the recommendations of the American Academy of Sleep Medicine (AASM) Manual for the Scoring of Sleep and Associated Events designed to guide sleep study professionals through the technical aspects of sleep testing.

Input signals used to derive outputs include:

- Saturation of Peripheral Oxygen (SpO₂)
- Photoplethysmogram (PPG)
- Pulse Rate

The autoscoring algorithm identifies the following physiological events based on AASM guidelines:

- Apnea-Hypopnea Index (eAHI)
- Respiratory Events (Apneas or Hypopneas): Obstructive, Central
- Sleep-Wake Events: Wake, Light Sleep, Deep Sleep, REM
- Total Sleep Time (TST)
- Total Recording Time
- Sleep Efficiency (SE)
- Sleep Latency (SL)
- Wake After Sleep Onset (WASO)
- Desaturation: Count, Index per Hour
- SpO₂ / Oxygen Saturation: Minimum, Maximum, Mean, Percent of Time in Range
- Heart Rate: Minimum, Maximum, Mean, Percent of Time in Range

Note: The scoring results should be reviewed and edited as necessary.

Clinical Performance

Device performance compared automatically detected events results obtained by a two-thirds majority of three experienced sleep technologists. Performance in terms of sensitivity and specificity was assessed at an AHI cut-off of $AHI \geq 5$. For more information about the clinical study, contact EnsoData or view the study information in the most recent device clearance.

The device was designed and tested under the following standards and guidelines:

- EN ISO 13485 Third Edition 2016/A11 Medical devices – Quality management systems – Requirements for regulatory purposes
- EN ISO 14971 Third Edition 2019-12 Medical devices - Application of risk management to medical devices
- IEC 62304 Edition 1.1 2015-06 Medical device software - Software life cycle processes

Apnea Hypopnea Index (eAHI)			
Desaturation	Sample Size (n)	Sensitivity	Specificity
3%	158	92.6% (87.2%, 97.2%)	71.6% (59.2%, 83.7%)
4%	158	89.4% (81.6%, 96.1%)	76.8% (67.1%, 85.4%)

Clinical trial performance in terms of sensitivity and specificity was assessed at an AHI cut-off of $AHI \geq 5$. The following table is supplemental information.

Apnea Hypopnea Index (eAHI) Supplemental Information				
Desaturation	eAHI Cutoff	Sample Size (n)	Sensitivity	Specificity
3%	15	158 (n=70 $AHI \geq 15$, n=88 $AHI \leq 15$)	74.1% (64.3%, 84.3%)	92.0% (86.4%, 96.6%)
4%		158 (n=40 $AHI \geq 15$, n=118 $AHI \leq 15$)	70.1% (55.0%, 82.6%)	95.8% (92.4%, 99.2%)
3%	30	158 (n=38 $AHI \geq 30$, n=120 $AHI \leq 30$)	53.0% (36.8%, 68.4%)	100% (100%, 100%)
4%		158 (n=13 $AHI \geq 30$, n=145 $AHI \leq 30$)	84.8% (61.5%, 100%)	98.6% (96.6%, 100%)

Sleep Staging			
Category	Sample Size (n)	Sensitivity	Specificity
Wake	52,622	86.7% (86.5%, 87.0%)	93.5% (93.4%, 93.7%)

Sleep Staging			
Category	Sample Size (n)	Sensitivity	Specificity
Light Non-REM	69,438	80.9% (80.6%, 81.2%)	85.5% (85.2%, 85.7%)
Deep Non-REM	10,195	63.4% (62.4%, 64.3%)	95.9% (95.7%, 96.0%)
REM	14,459	83.6% (83.0%, 84.2%)	97.5% (97.4%, 97.5%)

Sleep Profile and Oxygen Saturation					
Category	Deming Regression		Bland-Altman		
	Slope β_1	Intercept β_0	Mean Difference (MD)	Upper Limit (ULO)	Lower Limit (LLO)
eAHI (3%) [events/hour]	0.936 (0.853, 1.033)	0.023 (-1.185, 1.122)	1.000 (0.630, 1.367)	14.575 (13.779, 15.363)	-12.574 (-13.371, -11.786)
eAHI (4%) [events/hour]	0.982 (0.903, 1.130)	1.219 (0.116, 1.985)	-1.039 (-1.326, -0.749)	9.307 (8.692, 9.931)	-11.386 (-12.001, -10.763)
Total Sleep Time [hours]	1.159 (1.035, 1.318)	-0.695 (-1.576, -0.005)	-0.093 (-0.132, -0.059)	1.145 (1.060, 1.216)	-1.330 (-1.414, -1.259)
Sleep Efficiency [hours/hours]	1.154 (1.031, 1.317)	-0.088 (-0.205, 0.003)	-0.011 (-0.017, -0.007)	0.163 (0.151, 0.173)	-0.185 (-0.198, -0.176)
Sleep Latency [hours]	1.114 (0.997, 1.290)	-0.023 (-0.185, 0.090)	-0.129 (-0.154, -0.089)	0.884 (0.831, 0.970)	-1.143 (-1.196, -1.057)
Wake After Sleep Onset [hours]	1.073 (0.938, 1.219)	-0.271 (-0.436, -0.121)	0.167 (0.140, 0.196)	1.131 (1.073, 1.193)	-0.797 (-0.855, -0.735)
Oxygen Desaturation Index [events/ hours]	0.962 (0.896, 1.056)	1.667 (0.330, 2.847)	-1.046 (-1.417, -0.677)	13.223 (12.426, 14.015)	-15.315 (-16.111, -14.522)

Safety - Warnings and Cautions

- EnsoSleep PPG is a prescription device used under supervision of a physician in the United States.
- The device is not for patients with unstable cardiopulmonary conditions, current opiate or other narcotic drug use, or using home supplemental oxygen therapy, non-OSA respiratory ventilatory support therapy, diaphragmatic pacing, or cardiac stimulation therapy devices.
- The device is intended for analysis of sleep data acquired from adult patients.
- Capnography data is not supported or displayed.
- The device neither interprets results nor suggests a diagnosis or recommendations for treatment.
- The device aims to identify physiological events according to the latest sleep scoring rules, procedures, and definitions under the current standard of care as recommended by the AASM Manual for the Scoring of Sleep and Associated Events, industry-standard published papers, and methods cleared for this device.
- The scoring provided by the device must be reviewed, edited as necessary, and approved by a qualified healthcare professional.
- Any preliminary or preexisting score reports should be regenerated after review and rescoring of studies to ensure any/all scoring changes are reflected in final reports.
- Notify EnsoData and the appropriate government agencies if an adverse event has occurred that is related to use of this product.

EnsoData

EnsoData, Inc. reserves the right to make changes to both this document and to the product it describes. Product specifications are subject to change without notice. Nothing contained within this document is intended as any offer, warranty, promise or contractual condition, and must not be taken as such.

This product is approved for sale and distribution in the United States of America (USA). Do not use, resell, or distribute this product outside of the USA.

EnsoSleep PPG, EnsoTherapy, and EnsoSleep are trademarks of EnsoData in the United States. Non-EnsoData product names may be trademarks of their respective owners.
Published in the United States of America.



Manufactured by:
EnsoData, Inc.
10 E Doty Street, Suite 449
Madison, WI USA 53703

Part Number: REF0009-A-ENG
Publication Date: February 2024
Software Version: 7.1+

Phone: 608-889-3676 (ENSO)
Email: support@ensodata.com