

SOMNO*check* micro CARDIO

Simple Assessment of Cardiovascular Risk During Sleep

In sleep medicine - In internal medicine - In preventive medicine





In sleep diagnostics we provide classic data about apnea, desaturation and arousals.

We also have answers to questions that have long been open.

Which patient most urgently needs treatment?

The Cardiac Risk Index (CRI) is the new severity parameter in sleep diagnostics.

It supplements the well-known Apnea Hypopnea Index (AHI) with the cardiovascular risk dimension. So for the first time in an early screening of Sleep-Disordered Breathing (SDB), it is possible to see whether a patient is at risk of developing cardiovascular disease in addition to or as a result of SBD.

Which patient is really at risk?

The very first sleep screening made with SOMNOcheck micro CARDIO identifies patients with elevated cardiovascular risk. Furthermore, special indication is given of two additional risk factors for heart failure: arrhythmia (atrial fibrillation) and Cheyne-Stokes breathing.

Which additional diagnostics lead to the right diagnosis?

The CARDIO fingerprint for each patient shows which secondary or concomitant diseases may lead to an increased cardiovascular risk. This information shows the way to the next diagnostic steps – in or outside sleep medicine.



Our innovative measurement method implemented in the diagnostic device SOMNOcheck micro CARDIO works quickly, simply and non-invasively. For the first time a sleep screening device can determine a new severity parameter called "Cardiac Risk Index" (CRI) during a night-time recording. The CRI supplies information about your patient's cardiovascular risk, existent cardiovascular diseases and potential diagnostic steps.

A complete picture from only one nighttime recording

Conventional methods have shed light on particular aspects of cardiovascular risk, but SOMNOcheck micro CARDIO's measurement method is the first to yield a complete clinical picture with a single nighttime measurement. The Cardiac Risk Index is based on measuring actual changes in the patient, as opposed to calculating probability from statistic based risk scores. The algorithm in SOMNOcheck micro CARDIO has been validated against the risk scores of ESC/ESH.*

SOMNOcheck micro CARDIO also determines classic parameters such as desaturation and arousals and recognizes Sleep-Disordered Breathing, another important risk factor for cardiovascular diseases.

The basis of the analysis is the pulse oximetric measurement of the pulse wave. Changes in blood vessels are likewise detectable in the Pulse Wave Analysis (PWA), as are the responsiveness of the autonomic nervous system and heart rhythm. The same sensor measures oxygen saturation, which provides information about breathing disorders such as sleep apnea or Cheyne-Stokes breathing. By means of the optional nasal cannula, the screening device also detects mild obstructions in the patient's upper airways.



Grote L, Sommermeyer D, Zou D, Eder, D, N, Hedner J.
 Oximeter-Based autonomic state indicator algorithm for cardiovascular risk assessment.
 Chest 2011; 139:253-259



Explain increased risk with CARDIO fingerprint

The patient's own manifestations of the individual parameters go into the CARDIO fingerprint. Each parameter indicates certain diseases. The fingerprint serves as a compass, pointing toward the next diagnostic step and suggesting simple prioritization of patients.

- Innovative algorithms assess CRI:
 - Pulse wave amplitude analyses
 - Pulse wave reflection time and pulse rate analysis
 - Oxygen saturation analysis
 - Combinined measurement parameters
- Extensive risk status assessment:
 - Indication of endothelial dysfunction
 - Indication of central arterial stiffness
 - Recognition of pattern of diminished responsiveness
 - Detection of arrhythmia (atrial fibrillation)

Irregular pulse Low basal RCRD** saturation Low Frequent pulsedesaturation ratevariability Periodic pulse wave symmetric variability desaturation Short pulse wave reflection time Place the card on top of the CARDIO fingerprint and the diagnostic compass will show you the way. By the way: The card fits the original report print-out.

^{**} RCRD = Reduced Chronotropic Response to Desaturation

Sleep Diagnostics / SOMNOcheck micro CARDIO

Window	Displayed values	Source
Analysis time insufficient If there is no signal from either pulse oximetry sensor or flow sensor for more than two hours.	Analysis time insufficient	Analysis of artefact-free time per signal
Cardiovascular Risk	Low / Moderate / High CRI	Pulse wave analysis
Risk of sleep disorders Shows if patient is at risk of sleep-related breathing disorder	Low / Moderate / High Traffic light display: green, yellow, red	Analysis of results
Check for arrhythmia (AFIB)	Info window appears	Pulse rate analysis
Overview of Respiratory Events Apnea/Hypopnea Index Obstructive Apnea/Hypopnea Index Central Apnea/Hypopnea Index	AHI RDI OAHI ORDI CAHI CRDI	Flow signal: AHI. If this signal contains artefacts, an RDI determined by pulse oximetry and PWA will be displayed.
Check for Cheyne Stokes Breathing	Info window appears	Saturation analysis
Overview of Autonomic Arousals Autonomous Arousal Index Respiratory Autonomous Arousal Index Respiratory Effort Related Arousal Index (autonom)	AAI AAI resp RERAs	Pulse oximetry signal Pulse oximetry signal Pulse oximetry signal and flow signal
Overview of Oxygen Saturation Desaturation index Average Minimum	Drops Average Min	Pulse oximetry signal
Other Snore Average pulse rate Duration of recording	Snore Pulse av. Rec. time	Flow signal Pulse oximetry signal Analysis of artefact-free time
Artefact-free recording time If one of the two signals appears for less than four hours (i. e. several artefacts), a window opens that shows how long which signal was artefact-free.	Flow Pulse	Analyse artefact-free time per signal
Erase Data – erases all stored data	To erase press button for 3 sec	
Next Calibration – shows date of next recommended		Internal clock

Accessories for SOMNOcheck micro CARDIO



- Set of 100 nasal cannula, 90 cm WM 94522
 - | Softtip sensor CARDIO with Minimed plug (right-angled) Size. M: WM 94586 (not shown), Size. L: WM 94585
- Wristband WM 94560
- Transport bag WM 94055
- Software SOMNO*lab*, now with SOMNO*check* micro edition WM 98500
- 6 USB cable WM 94524
- Instructions for use SOMNOcheck micro CARDIO EN WM 96621 (not shown)
- Patient instructions for use SOMNOcheck micro CARDIO EN WM 96631 (not shown)

Our complete offering of therapy solutions accessories, mask systems and other technical data are at: **weinmann.de**

Software system requirements				
For a trouble-free installation of SOMNO <i>lab</i> , you will need administrator's rights on an IBM-compatible PC which fulfills the following requirements:				
Processor:	Pentium IV with 1,8 Ghz			
Available space:	Hard drive with at least 1 GB available memory and 1 GB available memory on a system partition			
Drive:	CD-ROM drive			
Input:	Keyboard and mouse or another pointer supported by Microsoft Windows			
Printer:	Supported by Microsoft Windows			
Operating system and main memory:				
	 Windows 2000 SP 4 or higher, if compatible with minimum 512 MB RAM, recommended 1024 MB RAM Windows XP 32 bit SP 2 or higher, if compatible with minimum 512 MB RAM, recommended 1024 MB RAM Windows 7 32 bit / Windows 7 64 bit with minimum 1024 MB RAM, recommended 2048 MB RAM Windows 8, 8.1, with minimum 1024 MB RAM, recommended 2048 MB RAM 			
Additional software:	- Internet Explorer 6.0 SP1 or higher if compatible - Adobe Acrobat Reader 6.0 or higher if compatible			

Software			
Data import via USB	Visualization of measurement data and events		
Event editing	Patient CARDIO fingerprint in report		
Extensive documentation Sleep-Disordered Breathing in report	Simple self-calibration – no maintenance costs		
Personalize the device	Programmable measurement time and duration		

Technical data SOMNO	check micro CARDIO	1	Certified Quality Management System meeting EC directive 93/42/EEC Annex II (EN ISO 13485)
Product class as per directive 93/42/EEC: Dimensions (W x H x D):	II a	■ Storage: -1	5°C to +40°C 0°C to +60°C 0°C to +60°C
Weight ■ Without batteries: ■ With batteries: Power supply:	79 g 145 g Type AA – Mignon	Pulse oximeter (Clipsensor) ■ SpO₂ measurement range: ■ SpO₂ accuracy 70 % < SpO₂ < 100 %: SpO₂ < 70 %: ■ Pulse rate measurement range:	45 to 100 % better than 2 % accuracy not validated 30 to 250 bpm
2 batteries (about	2 batteries (about 15 hrs.) 2 NiMH rechargeable batteries (about 20 hrs.)	■ Pulse accuracy:	1 bpm to 2 % of displayed value

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