



Innovative pulse oximeter with continuous monitoring

MySign® S from EnviteC is the intelligent solution for non-invasive vital sign monitoring.

- The sophisticated design guarantees ease of use, outstanding robustness and exceptional performance
- Versatile use in clinics, nursing homes, outpatient settings, emergency medical services and rescue situations
- Hygiene safety without compromise MySign® S is well protected, fast and easy to clean



Technical specification

Measuring range	Saturation (SpO ₂) 1 – 100% Pulse rate (PR) 0 – 300 BPM Perfusion index (PI) 0.1 – 20
Accuracy*	Saturation +/- 2% (70-100%, without motion artifacts) Pulse frequency +/- 3 BPM (30-250 BPM, without motion artifacts)
Display	Data update rate: once every 300 ms Plethysmogram: amplitude = 75% of the measuring range
Operating temperature	0°C-40°C (32°F-104°F)
Operating humidity	15-95% relative humidity (non-condensing)
Ambient pressure	700-1060 hPa
Storage temperature	-25°C-70°C (-13°F-158°F)
Battery	Li-ion 3.6 V 2900 mAh
Operating time per charge	> 18 hours (at standard settings)> 24 hours (when auto-off function is switched on and at lowest brightness setting for display)
Charger	Mini-USB Type B, Protection Class II Input: AC 110 V-230 V / 50-60 Hz / 125 mA, Output: DC 5 V / min. 1 A / < 15 W IEC 60601-1 / IEC 60950-1

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IP 54 IK 05 MDD 93/42/EWG DIN EN 1789





Approx. 4 hours Full functionality maintained 2.8" multicolor TFT 160 x 72 x 39 mm / 61/4 x 23/4 x 11/2 inch (H x W x D)
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IP 54
IK 05
Approx. 320 g (with sensor*)
USB 2.0 (Mini-USB Type B connector)
Monitoring of alarm limits and device functions (optical and acoustic)
Adjustable between: SpO ₂ Upper limit: 51%-100% SpO ₂ Lower limit: 50%-99% SpO ₂ Pulse Upper limit: 31-250 BPM Lower limit: 30-249 BPM
55-75 dB(A)
Tone pitch variable (depending on measured SpO ₂ value)
Period 0-8 hours
Max. 96 hours at a trend storage interval of 10 seconds (trend storage interval adjustable from 5-60 seconds) readings, date, time, alarm limits, events
Device and data set in connection with MySign® software (e.g. name, ward, patient ID)
Class II, Type BF
IIb
CE 0123
The device complies with the requirements of MDD 93/42/EEC for medical devices and the applicable standards (e.g. ISO 80601-2-61) and directives. Also in compliance with: DIN EN 1789 Medical vehicles and their equipment – Road ambulances *with SoftTin® R-3211-12 MySign®

*with SoftTip® R-3211-12 MySign®

All specifications apply to standard conditions: Ambient pressure 1013 hPa, 25°C dry ambient air.

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