



MARIE™ Capnograph

The Future of Respiratory Monitoring

Respiratory complications are among the most preventable, yet most serious threats to patient safety. Hypoventilation, CO₂ retention and apnea events often go undetected across many care settings, as continuous and reliable respiratory monitoring has traditionally been limited to intensive care and intubated patients.

EtCO₂

End-Tidal
Carbon Dioxide

RR

Respiration Rate

O₂

Supplementary
Oxygen Flow Rate

MARIE – a new standard in respiratory monitoring

MARIE is a wearable capnograph for non-intubated patients enabling advanced respiratory monitoring across care settings, from emergency to the ward.

Why MARIE?

- › Solves real problems: Detects respiratory deterioration, even during oxygen therapy
- › Expands access: Enables capnography and monitored oxygen therapy beyond the ICU
- › Designed for clinical use: Comfortable for patients and easy to use for staff

MARIE™ Capnograph

– Key Benefits

Wearable Capnography

Continuous EtCO₂, RR and O₂ flow display with clear waveform and trends.

Integrated Oxygen Delivery

Monitors and delivers supplementary oxygen quietly and comfortably up to 12 L/min.

Durable

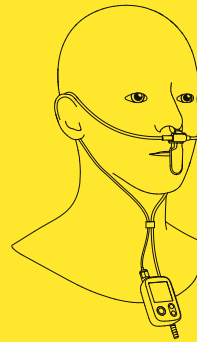
Up to 16 hours of battery life in a compact and rugged form factor.

Hassle-Free

No routine calibration required.

Applications

- › EMS & Ambulance Care
- › Emergency Departments
- › Procedural Sedation
- › Postoperative Monitoring
- › Wards & Step-Down Units



Essential Alerts

- › No Breath
- › Abnormal RR
- › Abnormal EtCO₂

Parameter	Specification
CO ₂	0–114 mmHg, 0–15 kPa / vol%
RR	3–150 breaths per minute
Oxygen Flowrate	0–12 L/min, up to approx. 60% FiO ₂
Trends	CO ₂ and Respiratory Rate, up to 8 hours selectable
Battery Life	Up to 16 hours, wireless charging
Weight	Monitor: 60 g, Sensor: 1.5 g
Alarms	No Breath, High/Low EtCO ₂ , High/Low RR, Rebreathing and technical alarms
Adapter Sets	Non-intubated
Patient Categories	Adults

Gas Measurements Accuracy According to EN/ISO 80601-2-55. The MARIE Capnograph is a CE-marked medical device (CE 2862) in accordance with Regulation (EU) 2017/745.