



Advantages of transcutaneous PCO₂ monitoring over PaCO₂ in chronic respiratory failure (CRF)

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Background

The study was aimed at assessing the accuracy of transcutaneous PCO₂ (PtcCO₂) monitoring compared to PaCO₂ and its technical drift during eight hours of nocturnal NIV in patients with stable chronic respiratory failure.

Methods

Only patients with a difference not greater than 2.5 mmHg between PaCO₂ drawn from the radial artery compared to PaCO₂ from the arterialized earlobe were included in the study. Accordingly, in 24 patients (15 with COPD) receiving nocturnal NIV capillary blood gases from the arterialized earlobe were analyzed at 11pm, 2am, 5am and 7am. Simultaneously, PtcCO₂ was monitored with the newest generation of three different devices: 1) SenTec Digital Monitor (SenTec DM), 2) TCM4-TINA, and 3) TOSCA500. A sensor temperature of 42°C was used for all devices, alarms were switched off.

Technical drift of the devices were analyzed after eight hours. In addition, method comparison to the gold standard PaCO₂ were performed, and analysis of patients' individual overnight CO₂-range were investigated in regard to intermittent measurements and continuous monitoring.

Table 1. Technical drift and continuous overnight PtcCO₂-monitoring. Median (interquartile range) is given for all patients (N=24).

	SenTec DM	TCM4-Tina	TOSCA500	p-value (RM-ANOVA)
Technical Drift (mmHg)	0.7 (-0.9/3.7)	-4.0 (-8.0/8.0)	-4.0 (-8.0/2.0)	<0.001*
Mean of overnight PtcCO ₂ (mmHg)	48.5 (36.5/65.9)	47.9 (31.5/68.6)	47.8 (37.6/65.9)	0.718
Standard deviation of overnight PtcCO ₂ (mmHg)	2.5 (1.1/11.5)	3.2 (0.3/23.0)	2.4 (1.5/12.1)	0.121

PtcCO₂ = transcutaneous partial pressure of carbon dioxide.

*lowest technical drift by SenTec DM compared to TCM4-Tina und TOSCA500.

Table 2. Method comparison of PtcCO₂/PaCO₂-pairs at 11 pm, 2 am, 5 am and 7 am. Results are given for all patients (N=24).

	SenTec DM	TCM4-Tina	TOSCA500
PaCO₂ versus drift-uncorrected PtcCO₂			
Pairs of values (N)	93	88	92
Mean of the difference (mmHg)	1.0	-1.5	0.8
Limits of agreement (range) (mmHg)	-4.7 to 6.7 (11.4)	-15.6 to 12.5 (28.1)	-6.8 to 8.3 (15.1)
PaCO₂ versus drift-corrected PtcCO₂			
	software based	calculated	calculated
Pairs of values (N)	90	80	92
Mean of the difference (mmHg)	0.8	-3.3	-1.6
Limits of agreement (range) (mmHg)	-4.9 to 6.5 (11.4)	-15.8 to 9.3 (25.1)	-8.4 to 5.2 (13.6)

PaCO₂ = arterial partial pressure of carbon dioxide, PtcCO₂ = transcutaneous partial pressure of carbon dioxide.

Figure 3. Overnight trend of PaCO₂ (red boxes) and PtcCO₂ (solid line represents SenTec DM, interrupted line represents TCM4-TINA, and dotted line represents TOSCA500) in a 34-year old patient with congenital central hypoventilation syndrome. Between 6.30am and 7am sensors of SenTec DM and TOSCA500 were disconnected from the patient.

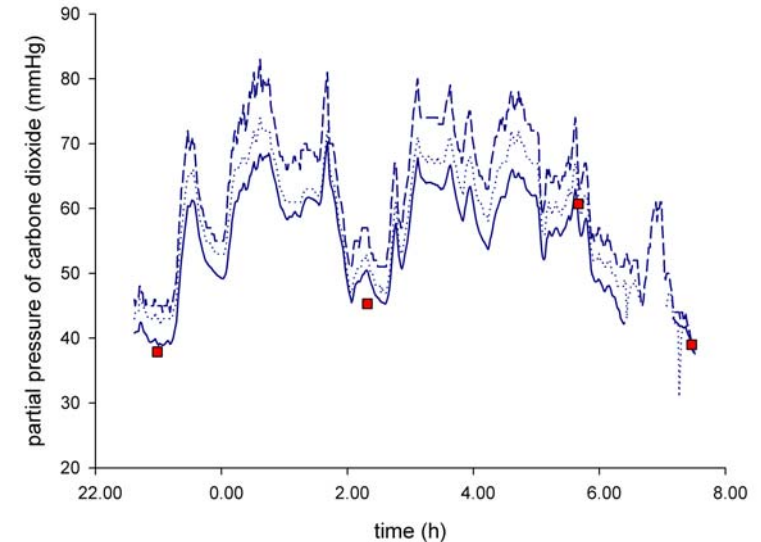


Figure 1: Range of overnight carbon dioxide measurements reflecting intermittent time spots at 11pm, 2am, 5am and 7am (RM-ANOVA, p=0.003).

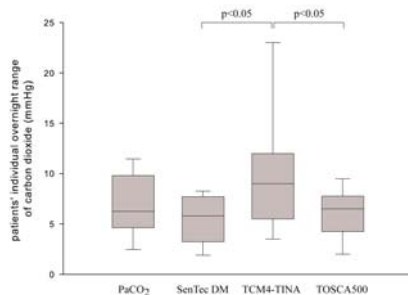
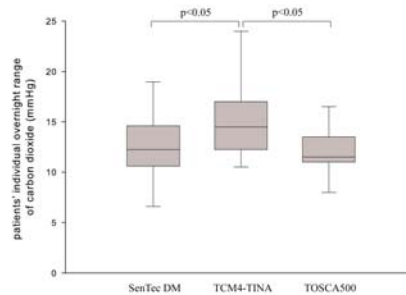


Figure 2: Range of overnight carbon dioxide measurements reflecting continuous transcutaneous monitoring (RM-ANOVA, p<0.001).



Results

The overall technical drift after 8 hours was appropriate for all devices regarding the lowest technical drift produced by SenTec DM, Table 1.

Method comparison by Bland-Altman of PaCO₂/PtcCO₂-pairs for drift-uncorrected and drift-corrected values are given in Table 2, respectively.

Individual overnight carbon dioxide ranges are illustrated for intermittent measurement points in Figure 1, and for continuous PtcCO₂-monitoring in Figure 2.

An overnight trend of PaCO₂ and PtcCO₂ in a 34-year old patient with congenital central hypoventilation syndrome is pictured in Figure 3.

Conclusion

As a result of technical refinements PtcCO₂-monitoring has now been shown to accurately reflect PaCO₂ in patients with nocturnal NIV. Since PtcCO₂-monitoring is also non-invasive, does not disrupt sleep quality and provides a more complete picture of alveolar ventilation than intermittent capillary PaCO₂, PtcCO₂ monitoring should become the preferred technique for assessing alveolar ventilation during nocturnal NIV.

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