



NAVSEA – Hypercapnia: Cognitive Effects and Monitoring

Year 2: Sensitivity, Reliability and Context Sensitive PO₂ Findings

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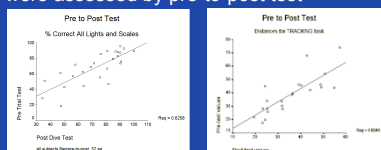
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Introduction:

This NAVSEA project is tasked to develop an algorithm to predict equivalent narcotic depth based on breathing gas inspired partial pressures. NASA's MATB-II flight simulator was chosen as the cognitive testing platform. The Multi-attribute task battery-II (MATB-II) is a suite of NASA-developed, computer-based, flight simulation software tests designed to evaluate operator performance and workload during operationally realistic scenarios. Here we present data from year 2 of the project showing: 1) pre-to-post trial score correlations 2) the protective effects of O₂ on CO₂ induced cognitive impairment at normobaric pressure and 3) the adverse effects of hyperbaric O₂ on CO₂ induced cognitive impairment in a scenario modeling CO₂ scrubber failure during a working dive.

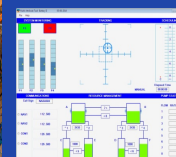
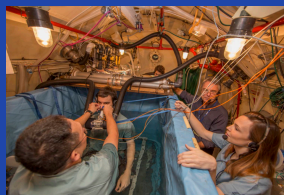
Methods:

After institutional approval and informed consent, as of 3/15/15, 28 subjects' baseline MATB-II scores (1 ATA air at rest) were compared to those recorded while breathing varying hyperbaric partial pressures of CO₂, N₂ and O₂ at rest and exercise. Inspired partial pressures tested were: (a) PiCO₂=0 and 0.075 ATA (7.5 kPa), (b) PiO₂=0.21 ATA (21.3 kPa), 1.0 ATA (101.3 kPa) and 1.22 ATA (123 kPa), and (c) PiN₂=0.79 ATA (80.0 kPa), 4.6 ATA (465 kPa) and 5.6 ATA (567.3 kPa). TRACKING scores were controlled for recent video game experience, exercise and pre-test hypercapnic ventilatory response (HCVR). The effects learning and breathing gas sequence were assessed by pre-to-post test correlations.



Results:

Arterial CO₂ and O₂ (for TRACKING task), and end-tidal CO₂, O₂, N₂ and their interactions (for SYSMON tasks) were significant predictors of performance by linear regression and repeated measures ANOVA respectively. The TRACKING and SCALES tasks were most sensitive. Pre-to-post test scores were well correlated (TRACKING Rsq= 0.59, SYSMON Rsq= 0.63, p<.01 two-tailed, Pearson). Preliminary data reveal a low to moderate narcosis signal for 4.5 to 5.6ATA N₂ and a strong narcosis / cognitive impairment signal for 0.075ATA CO₂. Quantifying the effect of O₂, however, is complicated in that it appears to affect cognition depending on its context.



Surface Oxygen effect: performance improved (Figure 1)

On the surface the study confirms the findings of Gill and Vann where O₂ appears to counteract CO₂ associated cognitive impairment. When compared to stages where subjects breathed either room air or 100% O₂, added CO₂ caused a 17 to 31% performance decrement both at rest and exercise (p<.001 ANOVA with Tukey's HSD post-hoc correction). However, post-hoc analysis showed significantly better performance during stages with higher PiO₂s where only a 7-12% decrement was observed.

Dive Oxygen effect: performance impaired (Figure 2)

During both normobaric and hyperbaric dives when PiN₂ was held constant at 4.5ATA a higher PiO₂ (1.2ATA) was associated with a 15 (normocarb) to 85% (hypercarb) performance decrement (p<.001) when compared to a normoxic PiO₂ (0.21ATA). Moreover, the higher PiO₂ (1.2ATA) was associated a 50% failure rate for completion of these test stages compared to normoxia. Six of the 12 exercise protocol subjects experienced potentially serious cognitive impairment while breathing added CO₂ with elevated PiO₂ and PiN₂ at depth. Affected subjects continued to fly the simulator, but entered a state of verbal unresponsiveness requiring physical intervention by the in-pool safety staff.

Elevated PiN₂ alone at 4.5 ATA did not significantly impair TRACKING performance (p=.366), however, additional data is forthcoming. Data from other MATB-II tasks (SYSMON, COMM and RESMAN) involving attention, memory and strategy show the expected N₂ narcosis effect.

Conclusions:

According to the Meyer-Overton theory of anesthetic potency, O₂ should cause cognitive impairment when breathed at hyperbaric pressures. Prior investigators have noted mixed effects; however, this may be because oxygen is a substrate for cellular respiration, a potent vasoconstrictor, and is CNS toxic at 1.6 ATA. Accordingly a partial pressure below 1.6 ATA may be insufficient to detect a measurable narcotic effect. Additional subjects and gas partial pressures should be tested to confirm these preliminary findings. Incipient CNS O₂ toxicity secondary to CO₂-mediated release of protective O₂ cerebral vasoconstriction will need to be distinguished from narcosis.

Figure 1 shows the mean TRACKING distances (± 95% CI) for immersed surface conditions. Performance impairment from 0.075ATA CO₂ was reversed by increased PiO₂= 925. (higher scores are worse)

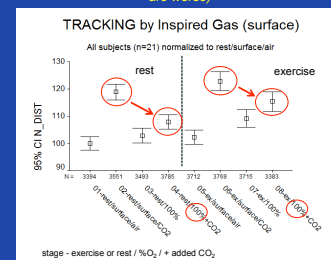
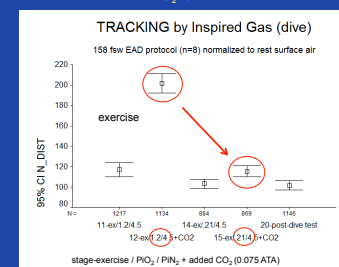


Figure 2 shows the severe impairment of TRACKING performance during 0.075ATA CO₂ challenge at 158 feet seawater while breathing a PiO₂=1.2ATA that was improved by switching to PiO₂=0.21ATA.



Gill M, Natoli MJ, Vacchiano C, MacLeod DB, Ikeda K, Qin M, Pollock NW, Moon RE, Pieper C, and Vann RD. Effects of elevated oxygen and carbon dioxide partial pressures on respiratory function and cognitive performance. Journal of applied physiology 117: 406-412, 2014.



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