

## Case report: Treatment of mild traumatic brain injury with hyperbaric oxygen

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### ABSTRACT

Two United States Air Force Airmen were injured in a roadside improvised explosive device (IED) blast in Iraq in January 2008. Both airmen suffered concussive injuries and developed irritability, sleep disturbances, headaches, memory difficulties and cognitive difficulties as symptoms of mild traumatic brain injury (mTBI). Six months after injury, repeat Automated Neuropsychological Assessment Metrics (ANAM) testing showed deterioration, when compared to pre-injury baseline ANAM assessment, in all measured areas (simple reaction time, procedural reaction time, code substitution learning, code substitution delayed, mathematical processing, and matching to sample).

The airmen were treated with hyperbaric oxygen in treatments of 100% oxygen for one hour at 1.5 atmospheres absolute, resulting in rapid improvement of headaches and sleep disturbances, improvement in all symptoms and resolution of most symptoms. Repeat ANAM testing after completion of the hyperbaric treatments — nine months after initial injury — showed improvement in all areas, with most measures improving to pre-injury baseline levels. The airmen received no other treatment besides medical monitoring. Repeat neuropsychologic testing confirmed the improvement. We conclude that the improvement in symptoms and ANAM performance is most likely attributable to HBO treatment.

### INTRODUCTION

Traumatic brain injury has been called one of the signature injuries of Operations Enduring Freedom and Iraqi Freedom. The RAND Report documented a 19% self-reported incidence of probable TBI among returning service members, with 320,000 probable TBI cases. Most of these cases (80%) are considered mild traumatic brain injury, or mTBI (1).

On a per-case basis, one-year costs for mTBI were estimated at \$27,259 to \$32,759 in 2007 (2). The lifetime costs of even mild TBI impairment in young service members can be deemed incalculable (3).

Mild TBI is usually characterized by a concussive event that causes a brief period of unconsciousness (lasting less than 30 minutes) or a period of confusion or amnesia lasting less than 24 hours. The Department of Defense has developed criteria for the diagnosis of mTBI, which must include one of the following:

- 1) any period of loss of or a decreased level of consciousness lasting less than 30 minutes;
- 2) any loss of memory for events immediately before or after the injury lasting less than 24 hours after the event;
- 3) any alteration in mental state at the time of the injury such as confusion, disorientation, or slowed thinking lasting less than 24 hours after the event;
- 4) transient neurological deficits (*e.g.*, weakness, loss of balance, change in vision, praxis, paresis or plegia, sensory loss, aphasia); and
- 5) normal intracranial imaging.

Findings may be transitory, and late sequelae that are not explainable by other means may qualify an individual for the diagnosis of mTBI. Patients with more than one of these findings may be assigned a higher level of TBI (4).



Since the symptoms of mTBI may develop gradually, are often subtle, and can be confused with other illness such as post traumatic stress disorder, mTBI may be unrecognized and undiagnosed (5). A concussive injury causes diffuse axonal injury, structural neuronal damage and diffuse neuronal dysfunction (6).

The symptoms of mTBI are variable and may include headache, irritability, impulsivity, anger, cognitive impairment, memory difficulty, loss of executive function, and vestibular and sleep disturbances (7). Electroencephalogram and sleep studies are usually normal. Most individuals with mTBI recover in three to 12 months, especially those who are young (8). However, some victims do not recover, or recover slowly; they are at risk for future injury and deterioration of brain function (9).

Mild TBI usually resolves without treatment within months, although approximately 20% of patients with mild TBI continue to have lingering symptoms for one year or longer after injury (1,10). Poor scores on neuropsychological testing months after injury have been correlated with poorer outcomes and unresolved symptoms (11).

Patients with several post-concussive symptoms are unlikely to improve after one year, in spite of traditional therapy (12). Treatment of mild TBI has included rest and observation, education, cognitive rehabilitation and pharmacotherapy (13).

Pharmacologic treatment may be required for control of disabling symptoms of headache, irritability, depression, and anger (14). Because of the efficacy of hyperbaric oxygen (HBO) in treating brain dysfunction from decompression sickness and carbon monoxide injuries, as well as anecdotal reports of its efficacy in treating concussive injuries, we felt HBO might prove of use in treating two airmen injured in a blast.

## **CASE REPORT**

In January 2008 Airman B, a 23-year-old male vehicle operator, was a convoy lead vehicle commander sitting in the passenger seat of an M915 14-ton truck. Airman C, a 22-year-old male vehicle operator, was driving the vehicle that was attacked with an improvised explosive device (IED).

The detonation occurred on the passenger side of the vehicle, nearer to where Airman B was sitting. The vehicle was damaged, and Airmen B and C sustained concussive injuries with a sense of being dazed for several minutes. There was no known direct blow to the head for either occupant or loss of consciousness, although both occupants had tinnitus. Airman B, who was approximately 3 feet closer to the blast, suffered immediately from a severe headache. Airman C continued to drive the damaged vehicle for several minutes and had no immediate symptoms other than being slightly dazed; however, he developed a mild headache some hours later.

Later in the day, Airmen B and C reported to the medical clinic, where no additional injuries were found. They were given acetaminophen for their headaches and placed on light duty. Two weeks later their symptoms had largely resolved, and they were returned to full duty.

Three weeks post-injury both airmen noted the return of headaches, with difficulty sleeping. Airman B expressed his headache severity as 5-6 and Airman C as 4-5 (on a scale of 1-10, with 10 being the most severe pain imaginable) with headaches occurring daily and lasting for several hours. Both individuals had difficulty falling and remaining asleep, and they reported sleep duration of three to six hours per night. Additionally both individuals felt they were quick to anger and stayed angry from trivial provocations for several hours. Lack of attention to detail, forgetfulness, and fatigue were also reported by both airmen. These latter symptoms began insidiously about three weeks after injury, progressed for about two months and remained constant for the next four months, until treatment with HBO was administered.

Upon arrival at their home base, the airmen presented to the clinic complaining of headaches, fatigue, lapses in memory, irritability and sleep disturbances. Neurological exams were normal, although the airmen appeared tired. Computerized tomography of the brain, EEGs and sleep studies were normal.

On initial deployment both airmen had received the Automated Neuropsychological Assessment Metrics test (ANAM) on 11 November



2007, two months prior to injury. This test was repeated on 21 July 2008, six months after injury. The repeat ANAM testing showed marked declines from the pre-injury baseline in several areas of measurement (*Figures 1A and 1B*, Page 394).

Airman B presented a statistically significant change in Simple Reaction Time and Matching to Sample tests, with declines in all other areas. Detailed neuropsychological testing of Airman B at six months post-injury and prior to HBO therapy revealed a diffuse or scattered pattern of deficits. Although his IQ score was within the average range, his neuropsychological functioning on a summary measure (Repeatable Battery for the Assessment of Neuropsychological Status – RBANS Form A) (15) was at just the 7th percentile.

Moreover, Airman B showed marked attention dysfunction for both auditory and visual material; cognitive processing speed was slowed and subjectively observed in casual conversation with the patient. He showed difficulty in repeating sentences and digit sequences as well as learning digit sequences over repeated trials.

Airman B also demonstrated problems in both verbal learning and visual memory. His reading speed was slowed, fingertip-tapping speed was slowed in both hands, and clerical speed for coding tasks was mildly impaired. He showed difficulty for rhythm perception and visual-motor integration for copying geometric designs. His reaction time was slowed on a computerized measure of attention. Reading level for sight words remained at the college level, but written arithmetic was at just the sixth-grade level.

Airman C presented statistically significant and drastic changes in both Simple Reaction Time modules (at the beginning and end of the battery), along with declines in all other areas except Mathematical Processing. Detailed neuropsychological testing of Airman C at the same time — prior to HBO therapy — was largely within normal limits notwithstanding problems for inconsistent attention and upper-right extremity dysfunction for grip strength and somatomotor integration. His RBANS (Form A) total score was at the 50th percentile, average range.

Initially, treatment of the headaches with ibuprofen and butalbital-aspirin-caffeine capsules (Fiorinal®) was tried, but these drugs were ineffective in relieving the pain. The airmen were placed on limited duty and daytime work only.

As the airmen had experienced at least one of the symptoms of mTBI after the blast (confusion, alteration of mental state) and their symptoms had no other reasonable explanation, they were given the diagnosis of mTBI in accordance with the Department of Defense criteria (4).

Because the two airmen had shown no improvement in their symptoms for seven months and were having difficulty performing their occupations, it was decided to begin hyperbaric oxygen treatment. Treatment with HBO was begun eight months post-initial injury. The treatment protocol was 100% oxygen for one hour at 1.5 atmospheres absolute. Treatments were given five days per week.

Clinical improvement was rapid. Airman C reported that his headaches vanished by the fifth treatment and did not return, and that he was able to sleep seven to eight hours per night uninterrupted. Airman B reported that his headaches weakened to 3-4 on a pain scale of 1-10, lasted only one to two hours instead of the previous eight to 10 hours, and that he was able to sleep eight to nine hours per night uninterrupted.

Both airmen reported that they felt more mentally alert and were less prone to forgetting, although they still did not feel “normal.” At the completion of the 40-treatment protocol, Airman C felt that his symptoms had ostensibly resolved, and Airman B felt that he was much improved, notwithstanding some lingering irritability and forgetfulness.

Repeat ANAM testing showed improvement in essentially all areas for both airmen. Airman C’s ANAM scores returned to pre-injury baseline levels, and Airman B’s ANAM scores returned to pre-injury levels, with no statistically significant differences in any of the tested domains (*Figures 1A and 1B*, Page 394, and *Figures 2A and 2B*, Pages 395-396).

Repeat detailed neuropsychological testing of Airman B showed improvement on some but not all

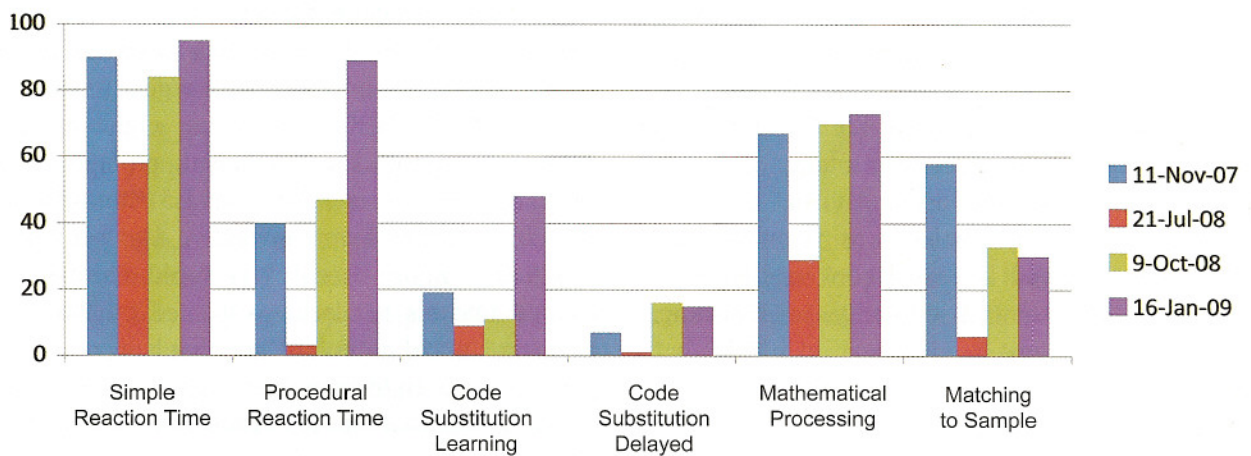


areas of cognitive functioning after HBO therapy at 10 months post-injury. His RBANS (Form B) total score was at the 12th percentile. For a patient with mild to moderate TBI, his scores improved faster than would be expected through spontaneous brain healing alone during this time interval. Areas of objective improvement included visuoconstructive abilities, fingertip-tapping speed and verbal learning/memory for word lists. His cognitive abilities status

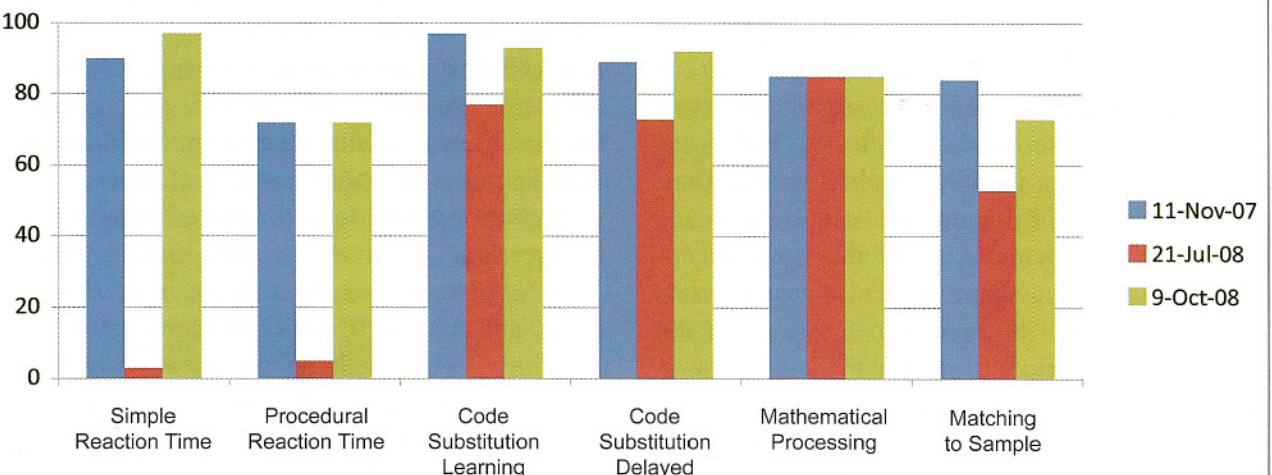
post-HBO treatment was deemed satisfactory to continue his job duties without special monitoring.

Repeat neuropsychological testing of Airman C was generally consistent with his pre-treatment test scores. Areas of subtle improvement such as motor abilities in the dominant right hand, written arithmetic and verbal fluency were observed. His RBANS B total score was at the 47th percentile, which was not a significant change from pre-treatment testing.

**FIGURE 1A — Airman B ANAM Scores**



**FIGURE 1B — Airman C ANAM Scores**



*Throughput scores are presented as the percentile of the comparison group of military members without TBI.*



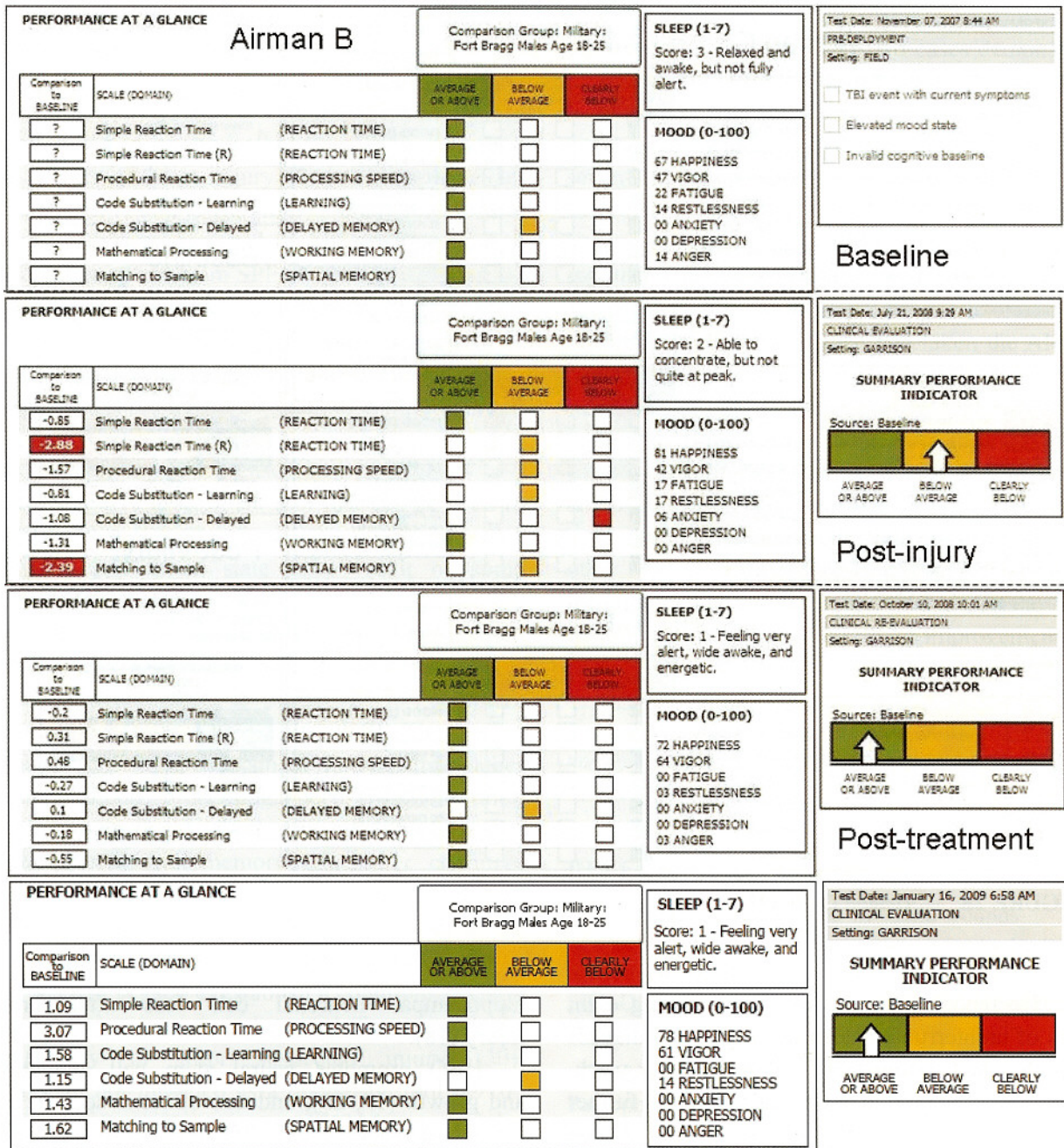


FIGURE 2A – Airman B ANAM Scores

Airman C was essentially well. Based on these results, it was decided to return Airman C to full duty, while Airman B continued hyperbaric treatment for another 40 treatments following the original treatment protocol.

Repeat ANAM testing on Airman B at the conclusion of the second set of 40 HBO treatments

showed improvement in all measures at or exceeding his pre-injury state, except for matching to sample, which was improved markedly from the injury state (Figures 1A and 2A, Page 394 and above).

Airman B reported that he had made continued improvement in cognitive function, felt much more alert and had returned to his pre-injury functional



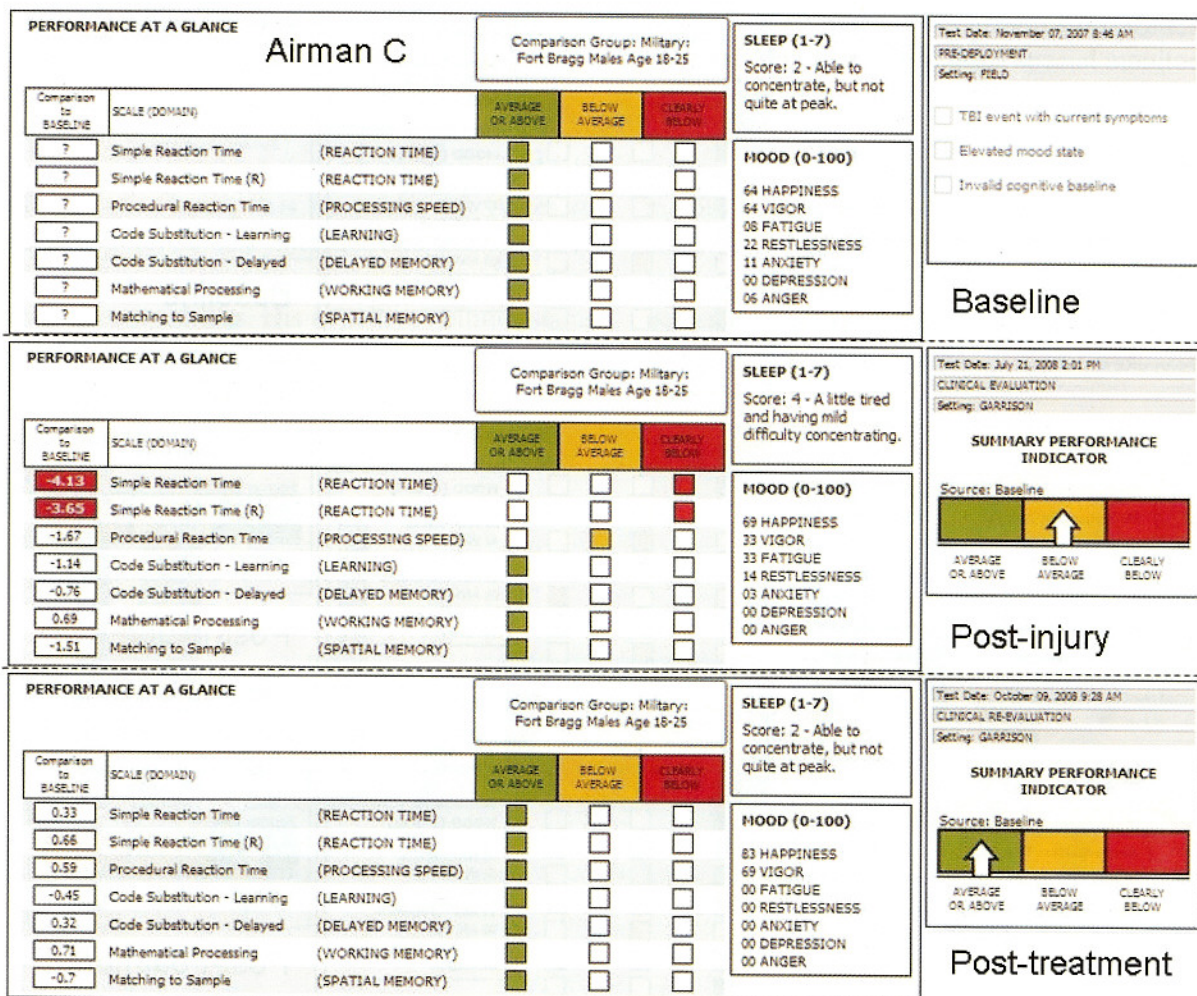


FIGURE 2B – Airman C ANAM Scores

state. He reported that he was experiencing eight hours of uninterrupted sleep per night, and that his headaches had diminished to about one per week.

He also noted the pain intensity had further decreased to 2-3 on a scale of 1-10, and that the headaches lasted two to three hours versus the original eight to 10 hours' duration.

**DISCUSSION**

Hyperbaric oxygen treatment has several effects that may be beneficial in treating brain injury. In animal models, HBO has been shown to enhance mitochondrial recovery and to reduce apoptosis in hypoxic nerve cells (16,17). The HBO-induced improvement in mitochondrial function appears to facilitate improved cognitive recovery and reduced

hippocampal neuronal cell loss after brain injury (18).

HBO promotes neural stem cell activation and growth (19, 20), and this effect is seen in the hypoxic-damaged brain (21). HBO also alleviates hypoxic-induced myelin damage, up-regulates HIF-1 alpha-enhancing neuronal tolerance to hypoxia, and increases cellular ATP levels and cognitive recovery after concussive injury (22).

Balance beam scores in rats with cerebral contusions were improved after treatment with HBO (23). In a rat model of chronic TBI, HBO improved spatial learning and increased vascular density in the injured hippocampus (24).

Controlled human studies of the efficacy of HBO after brain injury have been few. In a study



of moderate and severe TBI using the Glasgow Coma Scale and Glasgow Outcome Scale as measures of efficacy, an HBO-treated patient showed improvement over controls (25). HBO has been shown to be clinically effective in mediating the effects of brain injury (26). While the exact mechanism is unknown, HBO is thought to restore neural pathways damaged in TBI with supporting evidence supplied from SPECT brain imaging (27).

ANAM is a library of more than 30 computer-based test modules designed for a wide variety of clinical and research applications and is the direct outgrowth of more than 20 years of computer-based test development across all service branches within the Department of Defense (28). ANAM4™ is a neurocognitive assessment tool that can be used to identify changes in a service member's cognitive function and mood state as a result of some debilitating event.

The ANAM4™ TBI-MIL test battery used in this case report has been tailored to provide an instrument that is sensitive to cognitive changes that often accompany mTBI. The battery consists of a set of assessment modules that gather data on mood, processing speed (reaction time), working memory, short-term memory, spatial pattern recognition/memory and other cognitive functions. The test is designed for repeated testing and provides reliable measures when used for retesting as a measure of TBI recovery (29).

ANAM is used to establish a cognitive function baseline that can then be used for surveillance post-injury or after suspected injury (30). Although not intended as a diagnostic tool *per se*, comparative performance on ANAM test modules can be helpful in confirming the diagnosis, as demonstrated in this case report. In cases with known head trauma, computer-based assessments should be supplemented with detailed neuropsychological tests tailored to the patient's presenting problems and to the specific referral question to be answered.

## CONCLUSIONS

Several aspects of these two cases demonstrate the efficacy of HBO for the airmen treated. Although both airmen had stable symptoms of mTBI/post-concussive syndrome that had not improved for seven months, substantive improvement was achieved within 10 days of HBO treatment. The headaches and sleep disturbances improved rapidly, while the irritability, cognitive defects and memory difficulties improved more slowly.

Fortunately, both airmen had taken the ANAM and presented objective demonstration of their deficits from TBI and their improvements after HBO treatment. Both airmen, who were injured by the same blast sitting side by side, had similar symptom complexes of TBI and improved at similar rates after initiation of HBO treatment. Neither airman had any other form of treatment for TBI. It seems unlikely to the authors that any explanation other than the HBO treatments can be offered for their improvements. ■

*The views in this article are those of the authors and do not reflect the official policy of the Department of the Air Force, the Department of Defense or the U.S. Government.*

## REFERENCES

1. Tanielian T, Jaycox LH. Invisible wounds of war. Psychological and cognitive injuries, their consequences, and services to assist recovery. RAND Center for Military Health Policy Research. 2008.
2. Alexander MP. Mild traumatic brain injury: Pathophysiology, natural history, and clinical management. *Neurology*. 1995; 45:1253–60.
3. Gamboa AM Jr, Holland GH, Tierney JP, Gibson DS. American Community Survey: earnings and employment for persons with traumatic brain injury. *NeuroRehabilitation*. 2006; 21:327–33.
4. Assistant Secretary of Defense for Health Affairs. Health Affairs Memorandum (October 1, 2007). Traumatic Brain Injury: Definition and Reporting.



5. Practice parameter: the management of concussion in sports (summary statement). Report of the Quality Standards Subcommittee. *Neurology*. 1997; 48:581–5.
6. Maxwell WL, Povlishok JT, Graham DL. A mechanistic analysis of nondisruptive axonal injury: a review. *J Neurotrauma*. 1997; 14:419–40.
7. Arciniegas DB, Anderson CA, Topkoff J, McAllister TW. Mild traumatic brain injury: a neuropsychiatric approach to diagnosis, evaluation, and treatment. *Neuropsych Dis Treatment*. 2005; 311–27.
8. Carroll LJ, Cassidy JD, Peloso PM, Borg J, von Holst H, Holm L, Paniak C, Pépin M. Prognosis for mild traumatic brain injury: results of the WHO Collaborating Centre Task Force on Mild Traumatic Brain Injury. *J Rehabil Med*. 2004; (43 Suppl):84–105.
9. Biasca N, Maxwell WL. Minor traumatic brain injury in sports: a review in order to prevent neurological sequelae. *Prog Brain Res*. 2007; 161:263–91.
10. Elgmark Andersson E, Emanuelson I, Bjorklund R, Stålhammar D. Mild traumatic brain injuries: the impact of early intervention on late sequelae — a randomized controlled trial. *Acta Neurochirurgica*. 2007; 149, 151–9.
11. Sigurdardottir S, Andelic N, Roe C, Schanke AK. Cognitive recovery and predictors of functional outcome 1 year after traumatic brain injury. *J Int Neuropsychol Soc*. 2009; 15:740–50. Epub 2009 Jul 14.
12. Elgmark Andersson E, Emanuelson I, Olsson M, Stålhammar D, Starmark JE. The new Swedish Post-Concussion Symptoms questionnaire: a measure of symptoms after mild traumatic brain injury and its concurrent validity and inter-rater reliability. *J Rehabil Med*. 2006; 38:26–31.
13. Comper P, Bisschop SM, Carnide M, Tricco A. A systematic review of treatments for mild traumatic brain injury. *Brain Inj*. 2005; 19:863–80.
14. Veterans Health Initiative: Traumatic Brain Injury-Independent Study Course, Washington, D.C.: Department of Veterans Affairs, 2004 (<http://www1.va.gov/vhi/docs/TBI.pdf>).
15. Randolph, C. Repeatable Battery for the Assessment of Neuropsychological Status, San Antonio, TX: The Psychological Corporation, 1998.
16. Rockswold SB, Rockswold GL, Defillo A. Hyperbaric oxygen in traumatic brain injury. *Neurol Res*. 2007; 29:162–72.
17. Palzur E, Zaaroor M, Vlodaysky E, Milman F, Soustiel JF. Neuroprotective effect of hyperbaric oxygen therapy in brain injury is mediated by preservation of mitochondrial membrane properties. *Brain Res*. 2008; 1221:126–33. Epub 2008 May 11.
18. Zhou Z, Daugherty WP, Sun D, Levasseur JE, Altememi N, Hamm RJ, Rockswold GL, Bullock MR. Protection of mitochondrial function and improvement in cognitive recovery in rats treated with hyperbaric oxygen following lateral fluid-percussion injury. *J Neurosurg*. 2007; 106:687–94.
19. Wang XL, Zhao YS, Yang YJ, Xie M, Yu XH. Therapeutic window of hyperbaric oxygen therapy for hypoxic-ischemic brain damage in newborn rats. *Brain Res*. 2008; 1222:87–94. Epub 2008 May 18.
20. Yang YJ, Wang XL, Yu XH, Wang X, Xie M, Liu CT. Hyperbaric oxygen induces endogenous neural stem cells to proliferate and differentiate in hypoxic-ischemic brain damage in neonatal rats. *Undersea Hyperb Med*. 2008; 35:113–29.
21. Wang XL, Yang YJ, Xie M, Yu XH, Liu CT, Wang X. Proliferation of neural stem cells correlates with Wnt-3 protein in hypoxic-ischemic neonate rats after hyperbaric oxygen therapy. *Neuroreport*. 2007; 18:1753–6.
22. Peng Z, Ren P, Kang Z, Du J, Lian Q, Liu Y, Zhang JH, Sun X. Up-regulated HIF-1 alpha is involved in the hypoxic tolerance induced by hyperbaric oxygen preconditioning. *Brain Res*. 2008; 1212:71–8. Epub 2008 Mar 27.
23. Tinianow CL, Tinianow TK, Wilcox M. Effects of hyperbaric oxygen on focal brain contusions. *Biomed Sci Instrum*. 2000; 36:275–81.
24. Harch PG, Kriedt C, Van Meter KW, Sutherland RJ. Hyperbaric oxygen therapy improves spatial learning and memory in a rat model of chronic traumatic brain injury. *Brain Res*. 2007; 1174:120–9.



25. Lin JW, Tsai JT, Lee LM, Lin CM, Hung CC, Hung KS, Chen WY, Wei L, Ko CP, Su YK, Chiu WT. Effect of hyperbaric oxygen on patients with traumatic brain injury. *Acta Neurochir Suppl.* 2008; 101:145–9.
26. Shi XY, Tang ZQ, Xiong B, Bao JX, Sun D, Zhang YQ, Yao Y. Cerebral perfusion SPECT imaging for assessment of the effect of hyperbaric oxygen therapy on patients with post brain injury neural status. *Chin J Traumatol.* 2003; 6:346–9.
27. Harch PG, Fogarty EF, Staab PK, Van Meter K. Low pressure hyperbaric oxygen therapy and SPECT brain imaging in the treatment of blast-induced chronic traumatic brain injury (post-concussion syndrome) and post traumatic stress disorder: a case report. *Cases J.* 2009 Jun 9; 2:6538.
28. Vincent AS, Bleiberg J, Yan S, Ivins B, Reeves DL, Schwab K, Gilliland K, Schlegel R, Warden D. Reference data from the Automated Neuropsychological Assessment Metrics for use in traumatic brain injury in an active duty military sample. *Mil Med.* 2008; 173:836-52.
29. Segalowitz SJ, Mahaney P, Santesso DL, MacGregor L, Dywan J, Willer B. Retest reliability in adolescents of a computerized neuropsychological battery used to assess recovery from concussion. *NeuroRehabilitation.* 2007; 22:243–51.
30. Cernich A, Reeves D, Sun W, Bleiberg J. Automated Neuropsychological Assessment Metrics Sports Medicine Battery. *Arch Clin Neuropsychol.* 2007; 22 Suppl 1:S101–14. Epub 2006 Nov 21.

