REPETITIVE BLAST EXPOSURE AND BRAIN INJURY

Numerous research studies provide evidence that repetitive blast exposures harm the brain

I. Introduction

Trauma from blast exposure has become an increasing component of modern warfare. Blast injuries result from the sudden onset of shock waves generated by an explosion. Blast overpressure, an occupational hazard, is a significant cause of brain injury. This paper will first review scientific evidence that repetitive blast exposure harms the brain. It will underscore the vital need for blast pressure monitoring in military and tactical law enforcement personnel. Next, it will describe a recently available technology that enables this surveillance. Finally, it will emphasize the far-reaching benefits of blast pressure monitoring in protecting the health of military and tactical law enforcement personnel.

II. EVIDENCE THAT REPETITIVE BLAST EXPOSURE HARMS THE BRAIN The signature injury of modern warfare is mild traumatic brain injury (mTBI). Blast exposure is the leading cause of mTBI for US forces deployed to Afghanistan and Iraq in Operation Enduring Freedom (OEF), Operation Iraqi Freedom (OIF), and Operation New Dawn (OND).¹ The estimated prevalence of mTBI among returning service members is quite high, ranging from 15.2% to 22.8%.²-⁴ Acute symptoms of mTBI include headaches, dizziness, fatigue, irritability, confusion, memory problems, and sleep disturbances. In some cases, blast exposure can harm the brain without producing any obvious acute symptoms. Blast-related symptoms can surface much later in such cases.⁵

Research provides evidence that *blast exposure damages the structure of the brain.* In one study, Yeoh and colleagues reported evidence that blast overpressure disrupts the blood-brain barrier in rats.⁶ The blood-brain barrier normally protects the brain from harmful chemicals and regulates the transport of molecules into and out of the brain. In a separate study, Meabon and colleagues found evidence that blast exposure in mice damages the blood-brain barrier and interferes with the brain's expression of a protein called tau.⁷ In a postmortem study of veterans exposed to blast and/or concussive injury, Goldstein and colleagues found evidence of a tau protein-linked degenerative brain disease known as chronic traumatic encephalopathy (CTE). Veterans' brain pathology was similar to that of athletes who had a history of repetitive concussive injury.⁸

Using a swine model of blast-induced TBI, Ahmed and colleagues found evidence that blast exposure damages brain cells, compromises the permeability of blood vessels, and causes brain inflammation.⁹

In a study of 134 veterans, Robinson and colleagues found that close-range blast exposure was associated with altered brain connectivity, even when the blast exposure did not result in concussion symptoms at the time.¹⁰ In a study of 52 OEF/OIF veterans, Bazarian and colleagues found evidence that the severity of posttraumatic stress disorder (PTSD) is related to the severity of combat stress and structural brain changes, but not to a clinical diagnosis of mTBI. They concluded that blast exposure may induce subclinical brain injury and contribute to the onset of PTSD in a combat environment.¹¹



Furthermore, many studies have provided evidence that *the effects of blast exposure on the brain are cumulative and long-lasting.* In a study of 27,169 U.S. Army Special Operations Command (USASOC) personnel, Kontos and colleagues found that those with a history of blast-related mTBI were at greater risk of reporting PTSD symptoms than those with no mTBI history.¹² In a survey study, Carr and colleagues reported that repeated low-level occupational exposure to blast was associated with symptomatology similar to concussion. The number and severity of symptoms increased with history of blast exposure, and symptoms interfered with daily function.¹³

Meabon and colleagues found that in combat veterans with blast-related mTBI, the number of blast exposures correlated with symptoms of dizziness, loss of balance, and poor coordination. In addition, they found that an increase in the number of blast exposures was associated with lower glucose metabolism in the part of the brain known as the cerebellum. In a study of 80 USASOC personnel, Kontos and colleagues reported evidence that a history of blast-related mTBI exacerbates the initial symptoms of a subsequent mTBI. Trotter and colleagues, in a study of 249 veterans, found evidence that blast exposure accelerates the brain's aging process by reducing the integrity of the brain's white matter tissue. In a recent article, Elder and colleagues reviewed substantial evidence that blast-related TBI is

Review of Evidence and a Call to Action

pathophysiologically distinct from non-blast TBI and that low-level blast has long-term effects on the brain.¹⁶

III. THE CRITICAL NEED FOR BLAST PRESSURE MONITORING

Scientific studies have provided overwhelming evidence that blast exposure damages the brain, with long-lasting, adverse effects. Every day, military and tactical law enforcement personnel are forced to suffer the consequences of blast exposure. What else do we need to know about the effects of blast exposure? How could such knowledge help us to protect the health of military and tactical law enforcement personnel?

Our knowledge of the effects of blast exposure is severely limited by a lack of blast exposure monitoring. To estimate blast exposure, studies typically rely on self-report and semi-structured interviews of military personnel. Without monitoring blast pressure in individuals and evaluating the relationship between blast pressure levels and acute/chronic injury, we will never fully understand the cumulative effects of blast exposure. Obtaining objective measurements of blast exposure is critical for developing effective prevention and treatment strategies.⁵

Following the introduction of film badges for personal radiation monitoring in the 1920s, it took approximately 40 years to understand how radiation exposure related to the development of cancers. The Measurement was the necessary first step in understanding the cumulative effects of radiation exposure. Likewise, measurement of blast overpressure is the crucial first step in understanding the cumulative effects of blast exposure.

With objective, accurate measurements of blast pressure exposure, we would be able to correlate blast pressure levels

with levels of acute/chronic injury in individuals. The safety data from these studies would guide many key efforts. For example, this research would help us to identify blast pressure levels that put individuals at risk for brain injury and to tailor medical treatment to individual patients. It would also guide efforts to reduce blast pressure exposure and to determine how quickly to re-deploy individual service members.

IV. A Proven Technology for Blast Pressure Monitoring

A recently available, proven technology exists to monitor individual dosing of blast exposure in training and operations. The Blast Gauge® System utilizes a breakthrough, wearable sensor technology to document and quantify individualized blast exposure. It is a lightweight, three-sensor set that is worn on the helmet, chest, and shoulder.²⁰ Data from combat training, obtained from the Blast Gauge® System, show that military personnel experience *high rates of repetitive overpressure exposures* and support the need for cumulative overpressure monitoring.²¹⁻²² Blast Gauge® technology opens up a new approach to blast surveillance.

V. CONCLUSION AND CALL TO ACTION

Blast exposure is an occupational hazard that injures the brain. If we fail to take steps to reduce exposures, it represents a significant liability for the VA System. Now that the technology is available to measure individualized blast exposure, we need to use it to acquire data that will help us to develop effective prevention and treatment strategies. Measuring individualized blast exposure will provide critical data that will enable us to protect the long-term health of our warfighters and tactical law enforcement personnel.

VI DECEDENCES

- 1. McKee AC, Robinson ME. Military-related traumatic brain injury and neurodegeneration. Alzheimers Dement. 2014;10(3):S242-S253.
- 2. Hoge CW, McGurk D, Thomas JL, Cox AL, Engel CC, Castro CA. Mild traumatic brain injury in U.S. Soldiers returning from Iraq. N Engl J Med. 2008;358(5):453-463.
- Terrio H, Brenner LA, Ivins BJ, et al. Traumatic brain injury screening: Preliminary findings in a US Army Brigade Combat Team. J Head Trauma Rehabil. 2009;24(1):14-23.
 Polusny MA, Kehle SM, Nelson NW, Erbes CR, Arbisi PA, Thuras P. Longitudinal effects of mild traumatic brain injury and posttraumatic stress disorder comorbidity on postdeployment outcomes in National Guard soldiers deployed to Iraq. Arch Gen Psychiatry. 2011;68(1):79-89.
- Institute of Medicine. Gulf War and Health, Volume 9: Long-Term Effects of Blast Exposures. Washington, DC: National Academies Press; 2014.
- 6. Yeoh S, Bell ED, Monson KL. Distribution of blood-brain barrier disruption in primary blast injury. *Ann Biomed Eng.* 2013;41(10):2206-2214.
- 7. Meabon JS, Huber BR, Cross DJ, et al. Repetitive blast exposure in mice and combat veterans causes persistent cerebellar dysfunction. Sci Transl Med. 2016;8(321):321ra6.
- 8. Goldstein LE, Fisher AM, Tagge CA, et al. Chronic traumatic encephalopathy in blast-exposed military veterans and a blast neurotrauma mouse model. Sci Transl Med. 2012;4(134):134ra60.
- Ahmed F, Gyorgy A, Kamnaksh A, et al. Time-dependent changes of protein biomarker levels in the cerebrospinal fluid after blast traumatic brain injury. Electrophoresis. 2012;33(24):3705-3711.
- Robinson ME, Lindemer ER, Fonda JR, Milberg WP, McGlinchey RE, Salat DH. Close-range blast exposure is associated with altered functional connectivity in Veterans independent of concussion symptoms at time of exposure. Hum Brain Mapp. 2015;36(3):911-922.
- 11. Bazarian JJ, Donnelly K, Peterson DR, Warner GC, Zhu T, Zhong J. The relation between posttraumatic stress disorder and mild traumatic brain injury acquired during Operations Enduring Freedom and Iraqi Freedom. J Head Trauma Rehabil. 2013;28(1):1-12.
- 12. Kontos AP, Kotwal RS, Elbin RJ, et al. Residual effects of combat-related mild traumatic brain injury. *J Neurotrauma*. 2013;30(8):680-686.
- 13. Carr W, Polejaeva E, Grome A, et al. Relation of repeated low-level blast exposure with symptomology similar to concussion. *J Head Trauma Rehabil.* 2015;30(1):47-55.
- 14. Kontos AP, Elbin RJ, Kotwal RS, et al. The effects of combat-related mild traumatic brain injury (mTBI): Does blast mTBI history matter?. *J Trauma Acute Care Surg.* 2015;79(4):S146-S151.

 15. Trotter BB, Robinson ME, Milberg WP, McGlinchey RE, Salat DH. Military blast exposure, ageing and white matter integrity. *Brain.* 2015;138(Pt 8):2278-2292.
- 16. Elder GA, Stone JR, Ahlers ST. (2014). Effects of low-level blast exposure on the nervous system: is there really a controversy?. Front Neurol. 2014;5:269.
- Flakus FN. Detecting and measuring ionizing radiation a short history. *IAEA Bulletin*. 1981;23(4):31-36.
- 18. Inkret WC, Meinhold CB, Tashner JC. A brief history of radiation protection standards. Los Alamos Sci. 1995;23:116-123.
- 9. Khare P, Nair P, Khare A, Singh V, Chatterjee R. The road to radiation protection: A rocky path. J Clin Diagn Res. 2014;8(12):ZE01-ZE04.
- 20. BlackBox Biometrics, Inc. The Blast Gauge® System. https://blastgauge.com. Accessed March 7, 2016.
- 21. Bailie JM, Ma AB, Gomez R, et al. Blast exposure from shoulder mounted rocket launchers. Poster presented at: The 2015 Military Health System Research Symposium; August 17-20, 2015; Ft. Lauderdale, FL.
- 22. Da Silva UO, Lee G, Kamimori GH, Cho EG, Dark HE. Overpressure exposure to artillery and mortar crew members. Poster presented at: The 2015 Military Health System Research Symposium; August 17-20, 2015; Ft. Lauderdale, FL.