

Long-studied protein could be a measure of traumatic brain injury

July 13 2020

Scientists at the Walter Reed Army Institute for Research (WRAIR) have recently demonstrated that cathepsin B, a well-studied protein important to brain development and function, can be used as biomarker, or indicator of severity, for traumatic brain injury.

Traumatic brain injury (TBI) or brain trauma results from blows to the head, leading to life-changing disruption of the brain and a cascade of long-term health conditions. A leading cause of disability and death worldwide, TBI may occur due to an open-skull injury, like a gunshot wound, a fall, or an <u>automobile accident</u>. Athletes, the elderly, children, and <u>military service members</u> are particularly vulnerable.

Biomarkers are a source of great interest to researchers due to their potential to dramatically improve both the diagnosis and categorization of severity of TBI. Furthermore, they have the potential to validate treatment strategies by indicating whether drugs have reached their proposed targets and achieved therapeutic benefits.

In their publication in the *Journal of Neurotrauma*, the researchers showed that levels of cathepsin B were increased in areas of the injured brain relevant to controlling the senses, language, memory and other critical executive functions. In <u>healthy cells</u>, cathepsin B has a range of roles, including helping to eliminate damaged cells, maintaining metabolic homeostasis, and degrading improperly produced proteins. When the level of cathepsin B is not tightly controlled, it is linked to inflammation and tissue death. This publication reports the first results



demonstrating the ability to use cathepsin B as a blood-based biomarker to capable of identifying TBI severity within different brain regions as well as cerebral spinal fluid.

"Biomarker tests that accurately reflect the extent and severity of injury can dramatically improve the standard of care, minimizing the need for resource-intensive diagnostics like CT or MRI scans in favor of more portable tests," said Dr. Angela Boutte, lead author and section chief of molecular biology and proteomics within the Brain Trauma Neuroprotection Branch at WRAIR. "This would allow for early, accurate detection of TBI, whether at the side of the road after an accident or, most importantly, on the battlefield to help guide medical decisions."

Future research is planned to further characterize the role of cathepsin B in TBI.

More information: Angela M. Boutté et al, Penetrating Traumatic Brain Injury Triggers Dysregulation of Cathepsin B Protein Levels Independent of Cysteine Protease Activity in Brain and Cerebral Spinal Fluid, *Journal of Neurotrauma* (2020). DOI: 10.1089/neu.2019.6537

Provided by Walter Reed Army Institute of Research

Citation: Long-studied protein could be a measure of traumatic brain injury (2020, July 13) retrieved 20 February 2023 from <u>https://medicalxpress.com/news/2020-07-long-studied-protein-traumatic-brain-injury.html</u>

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