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Convergence insufficiency as a predictor of poor prognosis after acute mild traumatic brain injury

Kavya Devani¹, Neera Kapoor² and Latha Ganti^{3,4*}

Abstract

Background Mild traumatic brain injury (mTBI) is becoming a more common emergency department (ED) presentation. Towards this end, many types of testing in the acute setting are being investigated. One of these is screening for convergence insufficiency (CI) symptoms. These are common problems reported by patients with mTBI, but such oculomotor testing is rarely performed in the ED.

Objective To assess the feasibility of convergence insufficiency screening in the ED and investigate whether CI is associated with adverse events such as post-concussive symptoms or hospital admission.

Methods Written informed consent was obtained from patients age 18 years or older who experienced a mild head injury from any mechanism resulting in an mTBI. Patients underwent screening for CI symptoms using a standardized instrument of 15 questions, known as the convergence insufficiency symptom survey (CISS), with responses based on the Likert scale. These data were correlated to outcomes of hospital admission, occurrence of post-concussive symptoms, and 30-day hospital re-admission.

Results A total of 116 patients were prospectively enrolled, of which 58 were male. The median age was 31 years, with a range of 18 to 95 years of age. The median CISS score was 13, with an interquartile range (IQR) of 6 to 21 and an overall range of 0 to 53. Females presented with a median CISS score of 14, which was higher compared to the male median score of 10. The higher the CISS score, the more likely the patient was to be admitted to the hospital ($p = 0.0378$), develop symptoms of post-concussive syndrome at 30-day follow up ($p = 0.0322$), and be readmitted within 30 days ($p = 0.0098$).

Conclusions Screening for CI symptoms using the CISS can be a solid adjunct in the evaluation of mTBI in the ED. The CISS is easy and fast to administer, and it is a useful tool to stratify patients in terms of who is at the highest risk of developing complications related to the mTBI.

Keywords Convergence insufficiency, Traumatic brain injury, Head injury, Post-concussive syndrome, Extracranial manifestations of TBI

*Correspondence:

Latha Ganti

latha_ganti@brown.edu; lganti@ocom.org

¹ Vista Ridge High School, Cedar Park, TX 78613, USA

² New York University's Grossman School of Medicine, New York, NY 10016, USA

³ Orlando College of Osteopathic Medicine, Winter, FL 34787, USA

⁴ Warren Alpert Medical School of Brown University, Providence, RI 02903, USA



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Introduction

In 2020, the CDC documented 214,110 hospitalizations related to traumatic brain injury (TBI), with an additional 69,473 TBI-related deaths reported in 2021 [1]. This breaks down to a daily average of 586 hospitalizations and 10 deaths attributed to TBI, underscoring the widespread and serious nature of this injury [1]. Traumatic brain injury can be categorized into non-penetrating and penetrating forms. Non-penetrating TBI occurs when the brain sustains damage from a blow, bump, or jolt to the head or body. Conversely, penetrating TBI involves physical harm to the brain by passing through the skull [2]. While individuals of all ages, races, and genders can experience TBI, certain groups face a higher risk of enduring long-lasting effects due to previous trauma or limited access to advanced healthcare, as reported by the CDC [1]. Falls, firearm-related incidents, motor vehicle accidents, and assaults are common causes of TBI, with falls alone accounting for nearly half of TBI-related hospitalizations [3]. Approximately three quarters of TBIs that occur each year are concussions or other forms of mild TBI [4].

TBI can manifest through physical, cognitive, and perceptual/sensory symptoms [3]. Physical symptoms encompass headaches, vision changes, seizures, fluids from the ears or nose, nausea, or any neurological manifestation [3, 4]. Cognitive symptoms include alterations in consciousness levels, confusion, changes in sleep patterns, or behavioral shifts [3]. Changes in perception and sensations refer to alterations in the ability to interpret and process senses such as balance, taste, vision, hearing, and emotion [3, 4]. These symptoms arise from factors such as swelling, tearing, bruising, bleeding, or any alterations in the brain. The primary effects of TBI include diffuse axonal injury, concussion, hematomas, and contusions, as well as skull fractures, chronic traumatic encephalopathy, or post-traumatic dementia [3].

TBI severity is typically categorized as mild, moderate and severe based on the Glasgow Coma Scale (GCS), which assigns a score ranging from 3 to 15. This scale evaluates eye response (score out of 4), verbal response (score out of 5), and motor response (score out of 6) [5]. A higher score indicates a less severe injury. Specifically, a GCS score of 3 to 8 denotes severe TBI, 9 to 12 indicates moderate TBI and 13 to 15 indicates mild TBI (mTBI) [5]. However, a recent study of over 2200 hundred patients who presented to the ED with a mTBI demonstrated that there is in fact, nothing “mild” about a mTBI, even with a GCS of 15, much less one that is 13 or 14 [6].

This current study focuses on mTBIs and how screening for convergence insufficiency (CI) symptoms may be an efficient tool to risk-stratify patients in an emergency department (ED) setting. CI affects the ability

of both eyes to work together to form a single image, resulting from damage or dysfunction of nerves that control the eye muscles [7]. Although usually a problem that manifests in childhood, CI can occur following a brain injury as well [8]. Common symptoms include visual fatigue, double vision, headaches, or trouble concentrating [9].

This study focuses on the connection between the prognosis as well as symptom development regarding mTBI and the presentation of CI symptoms at the acute stage of mTBI at the ED. Since CI is a common sequela of mTBI, the goal of this study is to understand if CI symptomatology could be used as a predictor of prognosis to risk-stratify mTBI patients presenting to the ED.

Methods

This is a prospective observational study of consecutive adult patients who presented to the ED of a level I trauma facility with a mild TBI defined as a Glasgow Coma Scale (GCS) of 13–15. The mTBI must have occurred within the prior 24 h. Written informed consent was obtained from the patients to perform oculomotor testing including screening for CI symptoms using the CISS, in addition to their routine ED care.

The CISS is a validated 15-question instrument based on a Likert scale to assess the likelihood of CI, with “never” being 0 points and “always” being 4 points Fig. 1 [10]. Greater symptomatology related to CI is associated with a CISS score of 21 points or higher for adults and 16 and higher for children [10].

Consent also included follow-up telephone visits at 7 (3- to 15-day interval, defined as “early follow-up”) and 30 days (30- to 45-day interval, defined as “late follow-up”). Telephone follow-up visits included scripted questionnaires to assess whether patients had any symptoms suggestive of post-concussion syndrome (PCS), including headache, vomiting, dizziness, tinnitus, sensitivity to light, sensitivity to noise, numbness or tingling, blurred vision, diplopia, flashing lights, drowsiness, fatigue or lethargy, sadness or depression, nervousness or irritability, difficulty concentrating or remembering, sleeping problems, as well as feeling “slowed down,” “in a fog” or “dazed.” An affirmative response to any of these questions was considered to indicate the presence of PCS [11].

Data were entered into REDCap, a secure data collection tool that meets HIPAA compliance standards [12]. Statistical analyses were performed in JMP 16.0 for the Mac (Cary, NC) [13]. This study was conducted within the TBI ADAPTER trial [14] and was approved by the medical school's Institutional Review Board. Results of concurrent neurocognitive testing performed have been previously reported [15].

Convergence Insufficiency Symptom Survey

Name _____ DATE __/__/__

Clinician instructions: Read the following subject instructions and then each item exactly as written. If subject responds with "yes" - please qualify with frequency choices.

Do not give examples.

Subject instructions: Please answer the following questions about how your eyes feel when reading or doing close work.

		Never	(not very often) Infrequently	Sometimes	Fairly often	Always
1.	Do your eyes feel tired when reading or doing close work?					
2.	Do your eyes feel uncomfortable when reading or doing close work?					
3.	Do you have headaches when reading or doing close work?					
4.	Do you feel sleepy when reading or doing close work?					
5.	Do you lose concentration when reading or doing close work?					
6.	Do you have trouble remembering what you have read?					
7.	Do you have double vision when reading or doing close work?					
8.	Do you see the words move, jump, swim or appear to float on the page when reading or doing close work?					
9.	Do you feel like you read slowly?					
10.	Do your eyes ever hurt when reading or doing close work?					
11.	Do your eyes ever feel sore when reading or doing close work?					
12.	Do you feel a "pulling" feeling around your eyes when reading or doing close work?					
13.	Do you notice the words blurring or coming in and out of focus when reading or doing close work?					
14.	Do you lose your place while reading or doing close work?					
15.	Do you have to re-read the same line of words when reading?					
		<u> </u> x 0	<u> </u> x 1	<u> </u> x 2	<u> </u> x 3	<u> </u> x 4

TOTAL SCORE _____

Fig. 1 Validated Convergence Insufficiency Symptom Survey [10]

Results

The cohort consisted of 116 patients with mTBI, age range 18–95 years, with a median of 30.5 years, and IQR

of 21 to 50. The data were evenly split, with 58 males and 58 females. The racial distribution was 75% White, 19% Black, 3% Hispanic, and 3% other. The IQR for the score

distribution was 6–21, with the majority of patients CISS score less than 21. Females showed a higher median CISS score of 14 while males had a median CISS score of 10 (Fig. 2). A total of 32 patients (28%) had a CISS score ≥ 21 .

Figure 3 depicts a multivariate model that demonstrates a higher CISS score correlates with having

post-concussive syndrome (PCS). Specifically, symptoms of headache ($P=0.00004$), post-traumatic amnesia ($P=0.00062$), disorientation ($P=0.00547$), and difference in thinking or “fog” ($P=0.00622$). A higher CISS score was not significantly correlated with post-mTBI vomiting or altered state of consciousness. This model was robust,

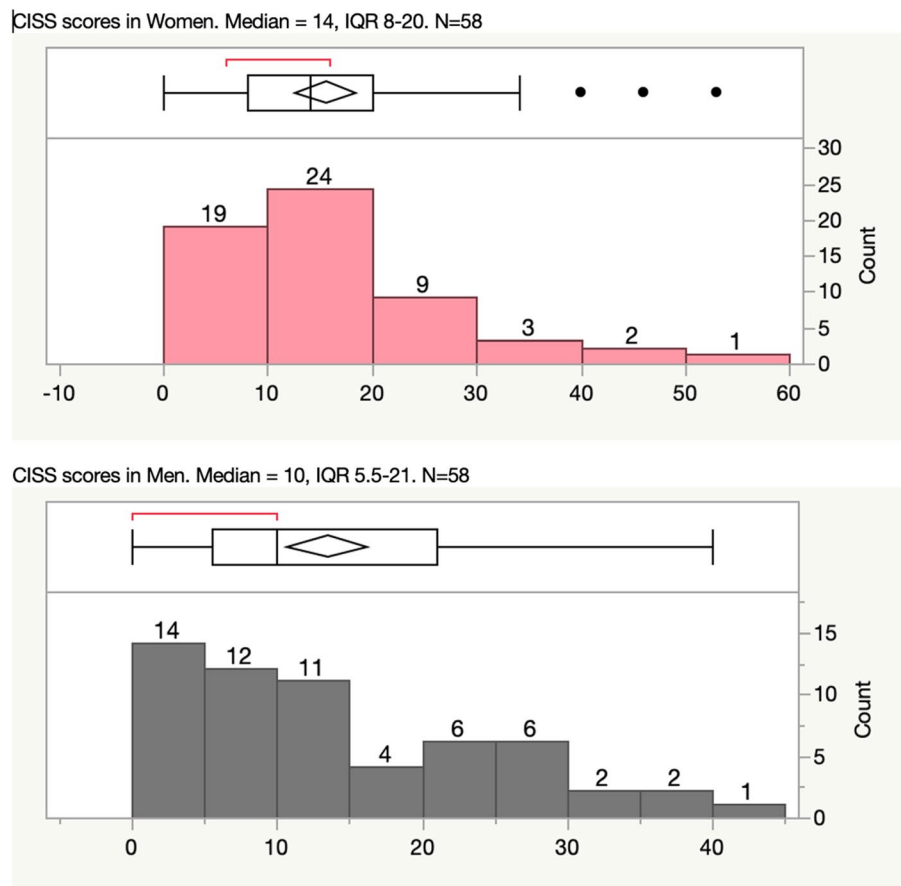


Fig. 2 CISS score distribution amongst men and women

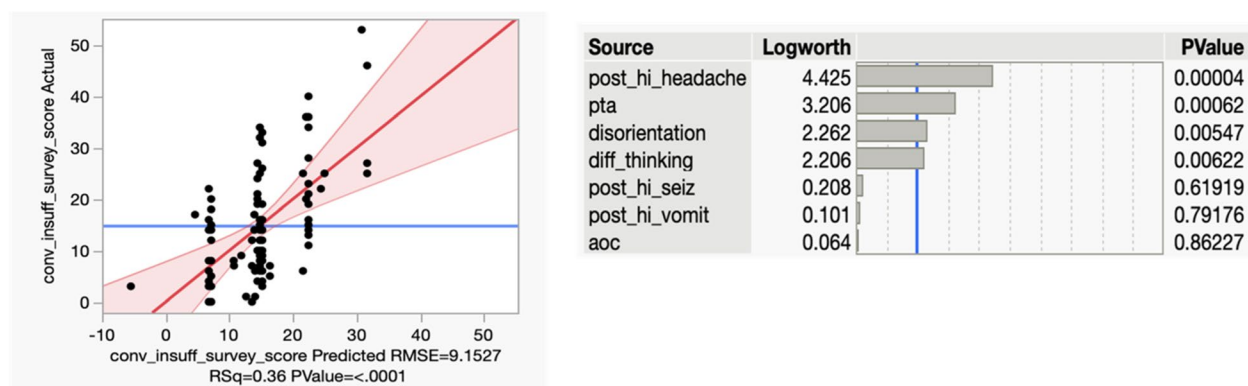


Fig. 3 Multivariate model depicting the correlation of CISS score to Post Concussive Syndrome after mild TBI

with an overall p-value of <0.0001 and a coefficient of determination (R^2) value of 36%.

A second multivariate model was built to analyze a potential correlation between the CISS score and the likelihood of being admitted for persisting symptoms related to mTBI Fig. 4. The model included age, sex, whether the initial computed tomography (CT) scan was abnormal, the ED's GCS score and the ED's CISS score. Figure 3 demonstrates that CISS is significantly correlated to hospital admission in this model, with a p-value of 0.02081. This model yielded an overall p-value of 0.0071, and a coefficient of determination (R^2) value of 17%.

A third multivariate model was built to analyze a potential correlation between the CISS score and the likelihood of being re-admitted within 30 days for persisting symptoms related to the mTBI Fig. 5. The model included age, sex, whether the initial computed tomography (CT) scan was abnormal, the ED's GCS score and the ED's CISS score. Figure 4 demonstrates that

CISS is significantly correlated to hospital admission in this model, with a p-value of 0.01248. This model was robust, with an overall p-value of 0.0118, and a coefficient of determination (R^2) value of 28%.

Discussion

This study aimed to assess the viability of administering the CISS in the ED and determine whether individuals over 18 years of age with a CISS greater than 21 were more prone to experiencing post-concussive symptoms and/or being admitted or re-admitted within 30 days. The results revealed a positive correlation between the CISS score and the development of post-concussive symptoms at a 30- to 45-day follow-up, as well as being admitted to the hospital at the initial ED visit and within 30 days thereafter. These data suggest that screening for CI symptoms with the CISS could be an effective tool for risk-stratifying patients, especially

Effect Summary

Variable	LogWorth	P-Value
Age	1.830	0.01478
CISS score	1.682	0.02081
CT scan abnormal?	0.764	0.17204
Sex	0.578	0.26397
ED GCS score	0.447	0.35697

ANOVA

Source	DF	Sum of Squares	Mean Square	F Ratio
Model	5	3.469412	0.693882	3.4295
Error	86	17.400153	0.202327	Prob > F
C. Total	91	20.869565		0.0071*

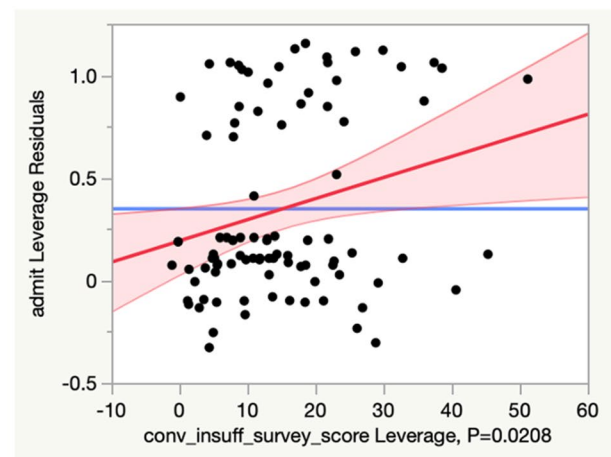
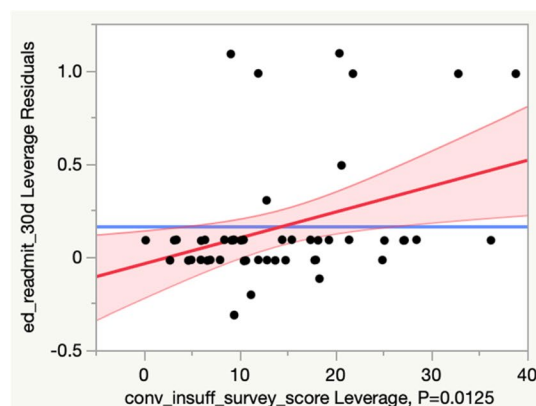


Fig. 4 Multivariate model depicting the correlation of CISS score to hospital admission after mild TBI



Source	LogWorth	P-Value
CISS score	1.904	0.01248
ED CT abnormal?	0.861	0.13779
ED GCS score	0.688	0.20490
Sex	0.427	0.37414
Age	0.002	0.99470

Fig. 5 Multivariate model depicting the correlation of CISS score to hospital re-admission within 30 days after mild TBI

given that the presentation of mTBI has become more prevalent in the ED setting.

A prior study of 72 TBI patients who were followed up at three years post-injury revealed a presence of vergence dysfunction in approximately 42% of these patients. In addition, these investigators reported that CI was associated with cognitive disturbances ($p < 0.005$), longer periods of coma ($p < 0.001$), and an inability of patients to find work in the open market ($p < 0.01$) [16]. Another study of 160 TBI patients reported that 90% presented with sensorimotor vision deficits, with accommodative and vergence deficits being the most common [17]. This suggests that CI portends a poor prognosis or further complications after a TBI. A review of visual impairments in the first year after traumatic brain injury encompassing 18 studies found that visual impairment negatively impacts independence in mobility and activities of daily living [18]. The investigators noted that the most common visual impairments seen include blurred vision, reading problems, diplopia, eyestrain, dizziness or disequilibrium in visually crowded environments, visual field defects, light sensitivity, and color blindness [19].

The significant association found between convergence insufficiency, hospital admission rates, and persisting post-concussive symptoms is biologically reasonable given the anatomy and physiology of extraocular motility. The nuclei for the three cranial nerves (oculomotor, trochlear, and abducens) innervating extraocular motility originate in the brain stem at the level of the midbrain. In addition to the midbrain and these three brain stem nuclei, other neurological areas that affect vergence include the frontal eye fields, mesencephalic reticular formation, medial longitudinal fasciculus, and cerebellum. Given these neuro-anatomical connections for vergence extraocular motility, it seems probable that mTBI might impact one or more of these areas impairing extraocular motility and possibly resulting in CI. Disruption to other neural tissues, vasculature, and cellular structures from an mTBI, beyond the discrete injury itself, lends credence to the fact that having a first concussion increases the probability of subsequent ones [20].

As TBI cases presenting to the ED continue to rise in numbers, it is of the utmost importance to identify factors that could help determine the need for early intervention. If supported by further analysis, screening for CI symptoms could guide triage for higher-level care and immediate intervention. It could serve as an identification tool for those at risk of post-concussive symptoms and other complications, offering care pathways to optimize both patient outcomes and healthcare utilization. Substantive work has been performed regarding vision rehabilitation of vision deficits following mTBI

[21–30], but larger-scale randomized controlled trials are indicated.

Conclusion

The current study demonstrates the feasibility of screening for CI symptoms in a busy ED setting for patients who experience mTBI or concussion. A higher CISS demonstrated a positive correlation with the development of PCS, being admitted to the hospital, and being at risk for re-admission within 30 days. Though more research is needed, this work offers a foundational idea to be built upon to target interventions toward at-risk mTBI patients in fast-paced, high-volume environments, such as the Emergency Department.

Authors' contributions

KD, NK, and LG participated in the design of the study. LG performed the statistical analysis. LG and NK conceived of the study and participated in its design and coordination. KD drafted the initial manuscript. NK and LG edited the manuscript. All authors read and approved the final manuscript.

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Data availability

No datasets were generated or analysed during the current study.

Declarations

Ethical approval and consent to participate

The study was performed in accordance with the ethical standards as laid down in the 1964 Declaration of Helsinki and its later amendments or comparable ethical standards. The University of Central Florida Institutional Board Review determined this study to be exempt.

Consent for publication

Not applicable.

Competing interests

Dr. Ganti has an editorial role at Springer.

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References

1. TBI data. <https://www.cdc.gov/traumaticbraininjury/data/index.html>. Accessed March 16, 2024.
2. Traumatic Brain Injury (TBI). <https://www.ninds.nih.gov/health-information/disorders/traumatic-brain-injury-tbi>. Accessed March 16, 2024.
3. Capizzi A, Woo J, Verduzco-Gutierrez M. Traumatic Brain Injury: an overview of Epidemiology, Pathophysiology, and Medical Management. *Med Clin North Am*. 2020;104(2):213–38. <https://doi.org/10.1016/j.mcna.2019.11.001>.
4. Stead LG, Bodhit AN, Patel PS, et al. Emergency Medicine Traumatic Brain Injury Research Network Investigators. TBI surveillance using the common data elements for traumatic brain injury: a population study. *Int J Emerg Med*. 2013;6(1):5. <https://doi.org/10.1186/1865-1380-6-5>.
5. Glasgow Coma Score. <https://www.ncbi.nlm.nih.gov/books/NBK513298/>. Accessed March 16, 2024.

6. Ganti L, Stead T, Daneshvar Y, Bodhit AN, Pulvino C, Ayala SW, Peters KR. GCS 15: when mild TBI isn't so mild. *Neurol Res Pract*. 2019;1:6. <https://doi.org/10.1186/s42466-018-0001-1>. PMID: 33324872; PMCID: PMC7650085.
7. Convergence Insufficiency. <https://www.nei.nih.gov/learn-about-eye-health/eye-conditions-and-diseases/convergence-insufficienc>. Accessed March 16, 2024.
8. Kapoor N, Balcer LJ, Rizzo JR. (2019) Vision problems. In *Textbook of Traumatic Brain Injury*, Third Edition (Edited by Silver JM, McAllister TW, and Arciniegas DB). American Psychiatric Publishing, Inc., Washington, DC, pp. 507–524.
9. Hajebrahami F, Gohel S, Scheiman M, Sangoi A, Iring-Sanchez S, Morales C, Santos EM, Alvarez TL. Altered large-scale Resting-State Functional Network Connectivity in Convergence Insufficiency Young adults compared with Binocularly Normal Controls. *Invest Ophthalmol Vis Sci*. 2023;64(14):29. <https://doi.org/10.1167/iov.64.14.29>. PMID: 37982763; PMCID: PMC10668612.
10. Rouse MW, Borsting EJ, Mitchell GL, Scheiman M, Cotter SA, Cooper J, Kulp MT, London R, Wensveen. Convergence Insufficiency Treatment Trial Group. Validity and reliability of the revised convergence insufficiency symptom survey in adults. *Ophthalmic Physiol Opt*. 2004;24(5):384–90. <https://doi.org/10.1111/j.1475-1313.2004.00202.x>. PMID: 15315652.
11. Ganti L, Khalid H, Patel PS, Daneshvar Y, Bodhit AN, Peters KR. Who gets post-concussion syndrome? An emergency department-based prospective analysis. *Int J Emerg Med*. 2014;7:31. <https://doi.org/10.1186/s12245-014-0031-6>. PMID: 25635191; PMCID: PMC4306054.
12. REDCap. <https://projectredcap.org/about/>. Accessed March 16, 2024.
13. JMP software. https://www.jmp.com/en_us/home.html. Accessed March 16, 2024.
14. Ganti L, Daneshvar Y, Bodhit A, et al. TBI ADAPTER: traumatic brain injury assessment diagnosis advocacy prevention and treatment from the emergency room—a prospective observational study. *Mil Med*. 2015;180(4):380–6. <https://doi.org/10.7205/MILMED-D-14-00316>.
15. Ganti L, Daneshvar Y, Ayala S, Bodhit AN, Peters KR. The value of neuro-cognitive testing for acute outcomes after mild traumatic brain injury. *Mil Med Res*. 2016;3:23. <https://doi.org/10.1186/s40779-016-0091-4>. PMID: 27453788; PMCID: PMC4957408.
16. Cohen M, Groswasser Z, Barchadski R, Appel A. Convergence insufficiency in brain-injured patients. *Brain Inj*. 1989;3(2):187–91. <https://doi.org/10.3109/02699058909004551>.
17. Ciuffreda KJ, Kapoor N, Rutner D, Suchoff IB, Han ME, Craig S. Occurrence of oculomotor dysfunctions in acquired brain injury: a retrospective analysis. *Optometry*. 2007;78(4):155–61. <https://doi.org/10.1016/j.optm.2006.11.011>. PMID: 17400136.
18. Greenwald BD, Kapoor N, Singh AD. Visual impairments in the first year after traumatic brain injury. *Brain Inj*. 2012;26(11):1338–59. <https://doi.org/10.3109/02699052.2012.706356>. Epub 2012 Aug 16. PMID: 22897509.
19. Blennow K, Hardy J, Zetterberg H. The neuropathology and neurobiology of traumatic brain injury. *Neuron*. 2012;76(5):886–99. <https://doi.org/10.1016/j.neuron.2012.11.021>. PMID: 23217738.
20. McCrea M, Guskiewicz K, Randolph C, Barr WB, Hammeke TA, Marshall SW, Kelly JP. Effects of a symptom-free waiting period on clinical outcome and risk of reinjury after sport-related concussion. *Neurosurgery*. 2009;65(5):876–82. <https://doi.org/10.1227/01.NEU.0000350155.89800.00>. discussion 882–3. PMID: 19834399.
21. Kapoor N, Ciuffreda KJ. Assessment of neuro-optometric rehabilitation using the Developmental Eye Movement (DEM) test in adults with acquired brain injury. *J Optom*. 2018;11(2):103–12. <https://doi.org/10.1016/j.optom.2017.01.001>. Epub 2017 Jul 1.
22. Ciuffreda KJ, Rutner D, Kapoor N, Suchoff IB, Craig S, Han ME. Vision therapy for oculomotor dysfunctions in acquired brain injury: a retrospective analysis. *Optometry*. 2008;79:18–22.
23. Thiagarajan P, Ciuffreda KJ. Effect of oculomotor rehabilitation on vergence responsivity in mild traumatic brain injury. *J Rehabil Res Dev*. 2013;50(9):1223–40.
24. Thiagarajan P, Ciuffreda KJ. Effect of oculomotor rehabilitation on accommodative responsivity in mild traumatic brain injury. *J Rehabil Res Dev*. 2014;51(2):175–91.
25. Thiagarajan P, Ciuffreda KJ. Versional eye tracking in mild traumatic brain injury (mTBI): effects of oculomotor training (OMT). *Brain Inj*. 2014;28(7):930–43.
26. Scharnweber AR, Palmer GA, Ampe HJ, Lenzen-Hammerel AM. Vision rehabilitation for traumatic brain injury and post-traumatic stress disorder. *Vis Dev Rehabilitation*. 2016;2(2):132–9.
27. Scheiman MM, Talasan H, Mitchell GL, Alvarez TL. Objective assessment of vergence after treatment of concussion-related CI: a pilot study. *Optom Vis Sci*. 2017;94(1):74–88.
28. Conrad JS, Mitchell GL, Kulp MT. Vision therapy for binocular dysfunction post brain injury. *Optom Vis Sci*. 2017;94(1):101–7. <https://doi.org/10.1097/OPX.0000000000000937>.
29. Ellis MJ, Leddy J, Cordingley D, Willer B. A physiological approach to assessment and rehabilitation of acute concussion in collegiate and professional athletes. *Front Neurol*. 2018;9:1115. <https://doi.org/10.3389/fneur.2018.01115>.
30. Fox SM, Koons P, Dang SH. Vision rehabilitation after traumatic brain injury. *Phys Med Rehabil Clin N Am*. 2019;30(1):171–88.

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