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**Dizziness Directly Influences Post-Concussion Symptoms and is Predictive of Poorer
Mental Health in UK Military Personnel: A Retrospective Analysis**

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ABSTRACT

Objective: To investigate the contribution of dizziness to post-concussion symptoms, depression, and anxiety symptoms. **Setting:** Mild Traumatic Brain Injury Service, Defence Medical Rehabilitation Centre, Stanford Hall. **Participants:** 283 UK military personnel from the Royal Navy, Royal Airforce, Royal Marines, and British Army. **Design:** A retrospective analysis of data from the Ministry of Defence (MoD) medical records database. **Main measures:** 16-item Rivermead Post Concussion Symptom Questionnaire, Generalized Anxiety Disorder 7-item scale, Patient Health Questionnaire-9, The Dizziness Handicap Inventory. **Results:** Injuries from sports or falls were the most common mechanism of mTBI (mild traumatic brain injury), accounting for 23% respectively. Chi-square analysis indicated that individuals with dizziness and post-concussion symptoms (PCS) had greater severity of PCS, depression, and anxiety than those with PCS alone. Mediation analysis showed dizziness directly and independently influenced the severity of PCS, despite the indirect effects of mediating depression and anxiety symptoms. **Conclusion:** Comorbid dizziness and PCS were predictive of poorer mental health compared with PCS alone. Additionally, dizziness directly influenced the severity of PCS irrespective of the indirect effects of mental health symptoms. These observations suggest that treating dizziness with vestibular rehabilitation may improve PCS and mental health.

Keywords: *Dizziness, mTBI, PCS, vestibular rehabilitation, UK military personnel.*

INTRODUCTION

During the conflicts in Iraq and Afghanistan, blast mild traumatic brain injury (mTBI) was characterized as a signature injury of the war and attributed to both increased exposure to explosive munitions, and higher survival rates due to advancements in battlefield medicine.¹⁻³ Currently, it is more likely that military personnel sustain blunt mTBI from civilian mechanisms such as sport, road traffic accidents (RTA), falls, and assaults whilst in garrison non-deployed settings, although published data to reflect this is sparse.⁴⁻⁵ What is well-known is that TBI is a global public health concern and a leading cause of death and disability, estimated to be sustained by 64 to 74 million civilians per year.⁶ Approximately 90% of these injuries are blunt-force and categorised as mild.⁴

Acute symptoms following mTBI typically resolve within three months, but 15% to 30% of individuals develop symptoms that persist for longer, sometimes years post-injury, known as post-concussion syndrome.⁷⁻⁹ This chronic condition is associated with a broad range of somatic, cognitive, and emotional symptoms and can be difficult to diagnose.¹⁰ This is partly because mTBIs frequently go unreported, but also because there are commonly no visible signs of anatomical damage from mTBI on computed tomography or magnetic resonance imaging scans when individuals do seek medical attention.¹¹⁻¹² Diagnosis of post-concussion syndrome is further complicated as the symptoms overlap with those from post-traumatic stress disorder (PTSD) and vestibular disorders.¹³⁻¹⁵

Dizziness is one of the most common symptoms of mTBI and vestibular disorders. Dizziness complaints have been shown to affect 84% of patients evaluated more than 30 days following blast exposure,¹⁶ with an enduring presence of postural instability evident up to 7 years post initial injury.¹⁷ However, vestibular pathology is not exclusive to blast mTBI and is frequently seen in blunt injuries too.¹⁸ These prevalences are especially concerning because dizziness at just 6 months post-onset is closely linked to psychological distress and a failure to return to work.¹⁹⁻²⁰

Different regions of the vestibular system are vulnerable to injury after both blunt and blast, head, or neck injuries. A retrospective study of 63 patients suffering from vertigo following TBI revealed several types of vestibular disorder²¹; benign paroxysmal positional vertigo (BPPV) was seen in 57% of these cases, cervicogenic vertigo in 27%, otolith disorder in 25%, labyrinth concussion in 19%, secondary endolymphatic hydrops in 19%, perilymphatic fistulae in 5%, and central vestibular in 5%. This variety of presentations highlights the need to carry out a full neuro-otological assessment to determine the most appropriate treatment.^{14,22-}

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Common difficulties that arise from mTBI, PTSD and vestibular dysfunction, all of which may occur simultaneously, encompass cognitive changes; such as memory and attention deficits, fatigue, anxiety, and depression.^{15,24-27} This similarity between symptom presentations may be partially attributable to overlapping pathophysiological changes such as neuroinflammation, excitotoxicity and oxidative damage.²⁸ There is a need to study these links further, to better understand the nature of mTBI and develop future treatment approaches.

Neuromodulation of the vestibular nerves in animal models also show that vestibular pathways to the hypothalamus are involved in stress responses mediated by the hypothalamic-pituitary-adrenal axis, which may well regulate function of ion transporters and ionic homeostasis of the inner ear.²⁹ This relationship is, however, bidirectional as more severe PTSD symptoms in US veterans were associated with worse dizziness handicap and vertigo symptom severity scores.³⁰ In fact, mediation analysis of UK military veterans with mTBI showed that vestibular disturbance directly and independently influenced increased severity of PCS, headaches, and disability; irrespective of mediating PTSD, depression, and anxiety.⁵ This, coupled with emerging evidence of the utility of vestibular rehabilitation therapies (VRT) in remediating PCS and PTSD as well as vestibular disorders, is indicative of the far-reaching influences of the vestibular system and the need to better understand them.³¹⁻³²

The main objective of the current study was to retrospectively examine data from UK military personnel with PCS seen via the mTBI service at the Defence Medical Rehabilitation Centre (DMRC) Stanford Hall. The aims were as follows: Firstly, to characterise the natural history of the mTBI sample. Secondly, to determine whether a combination of PCS and dizziness was predictive of poorer mental health outcomes compared with PCS alone. Thirdly, to evaluate whether dizziness directly influences the severity of PCS, independent of mediating comorbid depression and anxiety. Lastly, to examine the potential utility of VRT in the remediation of PCS.

METHODS

Participants

This study retrospectively reviewed the MoD internal medical records database of serving military personnel, including the Royal Navy, Royal Airforce, Royal Marines, and British Army. The sample comprised patients who had been referred to the mTBI service at the Defence Medical Rehabilitation Centre Stanford Hall for assessment and treatment of PCS. Only data from patients classified as having 1 or more mTBI were included, those with moderate or severe TBI were excluded. mTBI was classified by a loss of consciousness (LoC) from 0 to 30 minutes and/or post traumatic amnesia (PTA) from 0 to 1 day and/or alteration of consciousness (AoC) up to 24hrs, with normal structural imaging.³³⁻³⁴ Acute mTBI is within three months of sustaining injury, more than three months is a chronic condition. A favourable ethical opinion was granted prior to data collection from the University of Kent.

Data

Data review took place between March 2020 and October 2021. Records were looked at for patients who had been through the service between January 2016 and June 2021. Diagnosis of mTBI was confirmed by a Neurological Rehabilitation Consultant, Psychologist, Occupational Therapist, and Neuro Physiotherapist via a semi-structured clinical interview at the patient's first appointment with the DMRC Stanford Hall mTBI service. Prior to the

COVID-19 pandemic, these interviews were predominantly face-to-face, but from April 2020 onwards they were carried out via video conferencing platforms. The interviews determined the mechanism and severity of the brain injury, and assessed PCS presentation as well as comorbid depression and anxiety. Patients were diagnosed with PCS using a combination of the 16-item Rivermead Post Concussion Symptom Questionnaire (RPQ)³⁵ and clinical interview. Although $n = 8$ RPQ scores were not recorded in a way that we could access, all participants were diagnosed with PCS.

Upon entry to the mTBI service, most patients were additionally assessed for symptoms of anxiety using the Generalized Anxiety Disorder 7-item scale (GAD-7)³⁶, and depression, using the Patient Health Questionnaire-9 (PHQ-9).³⁷ The GAD-7³⁶ (anxiety) and PHQ-9³⁷ (depression) scales are 7-item and 9-item scales respectively, which ask participants to rate how often over the past two weeks they have experienced each symptom (item). Total possible scores range from 0-21 for the GAD-7 and 0-27 for the PHQ-9, with higher scores indicating more severe anxiety/depression.

Patients referred to the mTBI service with symptoms of dizziness and imbalance were also assessed to determine the severity of disability from dizziness using the Dizziness Handicap Inventory (DHI).³⁸ Where appropriate, patients' neuro-otological history was then assessed on an individual basis by a specialist neuro-physiotherapist, to determine the correct course and management of the specific balance and dizziness disorder.

Treatment

Following assessment, those patients who were considered to meet the clinical criteria for mTBI and who had ongoing symptoms were provided with education and support for managing their difficulties by a multi-disciplinary team encompassing neuropsychology, occupational therapy, and physiotherapy. Patients followed a tailored rehabilitation plan specific to their goals and symptoms, the aim being to help them manage any physical changes, cognitive difficulties, and psychological distress caused by the injury.

Treatment of dizziness symptoms consisted predominantly of a VRT program, which among other components encompassed gait training, balance and habituation exercises, gaze stability work and breathing techniques, and canalith repositioning manoeuvres (e.g., barbecue roll, Epley) for benign paroxysmal positional vertigo (BPPV). The rehabilitation exercises were performed within a range of indoor (e.g., gym, pool) and outdoor settings, as well as within a virtual reality environment, and BPPV treatment was repeated as often as necessary alongside other mTBI treatment. Patients who required more intensive input for persistent symptoms (vestibular and otherwise) were additionally invited to attend a group residential course which targeted areas such as fatigue management, relaxation training, and graded return to exercise.

Statistical Analysis

Summary statistics were calculated to show sample demographics, mTBI history, and comorbid symptoms. Two chi-square analyses were then performed to determine the relative frequency and severity of depression (PHQ-9) and anxiety (GAD-7). This included two groups: those with PCS and dizziness, and those with PCS only. Individuals suffering from dizziness and PCS were diagnosed by clinicians via a combination of objective (e.g., physical examination tests) and subjective (e.g., symptom questionnaire and clinical interview) assessments. Independent samples *t* tests were used to investigate differences in the severity of comorbid symptoms between those with and without dizziness, and a mixed Analysis of Variance (ANOVA) was then performed on the two groups to compare differences in PCS (as measured by the RPQ) pre- and post- treatment. Finally, in the PCS-dizziness group a mediation analysis was implemented to establish whether the severity of dizziness (as measured by the DHI) directly influenced PCS (RPQ) when depression (PHQ-9) and anxiety (GAD-7) were taken into account as mediators. This analysis also examined the combined total effects of dizziness in the mediation and outcome variables. All analyses were conducted using SPSS 26 and participants with missing data were excluded. Mediation analysis was conducted

utilising Hayes³⁹ macro for SPSS with bias correction bootstrapping the sample to 10,000 with 95% confidence intervals. Coefficients were considered statistically significant at $P < .05$.

RESULTS

Overview of Sample Characteristics

As can be seen from the demographics in Table 1, the majority of the sample were male (86%) non-commissioned officers (89%), who predominantly served in the British Army (72%). Ages ranged from 18-62, with mean age 32.2 (SD 8.8).

Table 1. *Sample demographics (N=283)*

		<i>n</i>			<i>n</i>
Gender	Male	243	Military service branch	Army	203
	Female	40		Airforce	42
Rank	Non-Commissioned officers	251		Navy	29
	Commissioned officers	32		Royal Marines	9

mTBI history can be seen in Table 2. The two most frequently reported mechanisms of injury were sports and falls, each equally accounting for 23% of the sample. More than half of the participants had suffered post-traumatic amnesia for less than 24 hours (60%) and/or a loss of consciousness for less than 30 minutes (53%). Altered consciousness was experienced by (39%) of the sample and 48% had a previous history of more than 1 mTBI. Two thirds of the sample had chronic PCS (65%) and 35% had acute PCS and were seen within three months of sustaining their injury.

Table 2. *Overview of mTBI history (N=283)*

		<i>n</i>			<i>n</i>	
Mechanism of Injury	Fall	64	mTBI History	>1 mTBI	135	
	Sports-related	64		Level of Consciousness	Loss of consciousness	149
	Road traffic accident	52			Altered consciousness	109

Assault	45	PCS Symptoms	Post-traumatic amnesia	169
Blast	28		Chronic	185
Other	30		Acute	98

Abbreviations: mTBI, mild traumatic brain injury; PCS, postconcussion symptoms

GAD-7, PHQ-9, DHI, and RPQ scores were captured during initial clinical assessment. As can be seen in Table 3, these showed that more than half of the participants (58%) reported symptoms of anxiety, two-thirds (67%) reported symptoms of depression, and 45% of the sample reported symptoms of dizziness. Overall, 60% of the sample were diagnosed with PCS only and 40% were diagnosed with dizziness and PCS.

Table 3: Frequency of co-morbid symptoms (N=283)

		Missing			
		<i>n</i>	cases (<i>n</i>)		
GAD-7^a	Anxiety present	165	36	Diagnosis/ PCS Group	
PHQ-9^a	Depression present	189	35	PCS only	171 0
DHI^b	Dizziness present	127	143	Dizziness/ PCS	112 0

Abbreviations: DHI, Dizziness Handicap Inventory; GAD-7, Generalized Anxiety Disorder 7-item scale; PCS, postconcussion symptoms; PHQ-9, Patient Health Questionnaire-9.

^aPresence scores of 5 or more³⁶⁻³⁷

^bPresence scores of more than 0

The Influence of Dizziness on GAD-7, PHQ-9, and RPQ Scores

Exploratory chi-square analyses indicated that there was a significant association between severity of depression (PHQ-9) in relation to PCS group, $\chi^2_2=15.759$, $P<.001$. Specifically, those with dizziness and PCS were more likely to have moderate to severe levels of depression than those with PCS only. There was also a significant association between PCS group and anxiety (GAD-7), $\chi^2_1=5.610$, $P=.018$; individuals with PCS only were more likely to have no or mild anxiety, whereas those with dizziness and PCS were more likely to have moderate to severe levels of anxiety.

The Effect of Dizziness on Comorbid Symptoms

As can be seen in Table 4, independent samples *t* tests revealed that the PCS-dizziness group had significantly worse symptoms of depression, dizziness, and PCS pre-treatment, compared with the PCS-only group.

Table 4. Comparison of comorbid symptom severity in those with and those without dizziness

Measure	Dizziness-PCS		PCS-only		<i>t</i> _{df}	<i>P</i>	Cohen's <i>d</i>
	Mean	SD	Mean	SD			
Depression (PHQ-9)	11.8	6.4	8.4	6.3	4.1 ₂₄₆	<.001 ^a	0.5
Anxiety (GAD-7)	9.2	5.8	7.5	5.8	2.3 ₂₄₅	.025	0.3
Dizziness (DHI)	36.9	23.3	18.7	19.1	4.3 ₁₃₈	<.001 ^a	0.9
Pre-treatment PCS (RPQ)	31.2	14.3	25.0	13.8	3.6 ₂₇₃	<.001 ^a	0.4
Post-treatment PCS (RPQ)	18.3	14.9	15.7	13.7	1.2 ₁₇₄	.232	0.2

Abbreviations: DHI, Dizziness Handicap Inventory; GAD-7, Generalized Anxiety Disorder 7-item scale; PCS, postconcussion symptoms;

PHQ-9, Patient Health Questionnaire-9; RPQ, Rivermead Post Concussion Symptoms Questionnaire.

^aSignificant when adjustments are made for multiple comparisons ($P=.05/5=.01$)

A Comparison of RPQ Scores Pre-and Post-Treatment

To compare PCS scores before and after treatment, a 2 (PCS group: PCS-dizziness vs. PCS-only) x 2 (Time: pre-treatment vs. post-treatment) mixed ANOVA was performed on RPQ scores. This showed a significant main effect of PCS group, $F_{1,170}=4.3$, $P=.040$, $\eta_p^2=.025$, observed power=0.5, with the PCS-dizziness group having significantly higher RPQ scores (mean 25.1, SE 1.3) than the PCS-only group (mean 21.3, SE 1.3). There was also a significant main effect of Time, $F_{1,170}=131.4$, $P<.001$, $\eta_p^2=.436$, observed power=1.0, with participants having significantly lower RPQ scores post-treatment (mean 17.2, SE 1.1) compared with before (mean 29.1, SE 1.0). There was no Group x Time interaction effect, $F_{1,170}=1.0$, $P=.317$, $\eta_p^2=.006$, observed power=0.2.

The Influence of Dizziness on RPQ Scores

Prior to mediation analysis a multiple linear regression was conducted to identify the variables that were significantly associated with the DHI scores. DHI, PHQ-9, GAD-7, and RPQ scores were all significantly associated with each other (all $P < .001$ with coefficient scores ranging from $r = 0.554$ to $r = 0.726$). A mediation analysis was then performed to determine whether the degree of dizziness seen in participants with dizziness and PCS before treatment had a direct effect on their pre-treatment RPQ scores and any indirect effects on their anxiety (GAD-7) and depression (PHQ-9) scores.

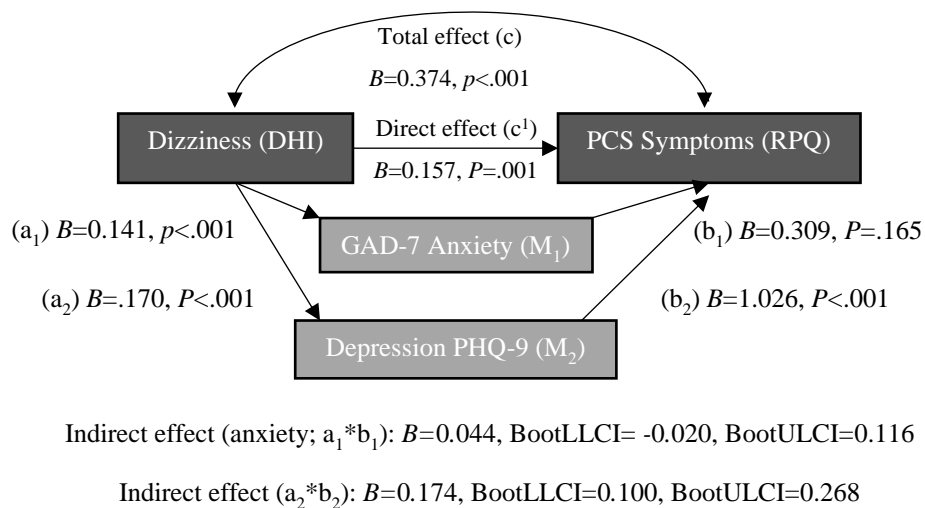


Figure 1. Mediation Analysis. RPQ ($N = 132$). DHI indicates Dizziness Handicap Inventory; GAD-7, Generalized Anxiety Disorder 7-item scale; PCS, postconcussion symptoms; PHQ-9, Patient Health Questionnaire-9; RPQ, Rivermead Post Concussion Symptoms Questionnaire.

As can be seen from Figure 1, there was a direct effect of dizziness on pre-treatment PCS scores, $P = .001$. A significant association in pathway a_1 was seen ($P < .001$), with dizziness influencing anxiety and depression in pathway a_2 ($P < .001$). Although there was no indirect effect of anxiety on PCS (pathway b_1 , $P = .165$), depression did have an indirect effect (pathway b_2 , $P < .001$). Overall, dizziness, anxiety and depression were shown to be significantly associated with the total effects of PCS ($P < .001$).

DISCUSSION

The main findings of this retrospective analysis indicate that dizziness both directly and in conjunction with the total effects of mental health influences the severity of PCS. The strength of these directional associations are supported by chi-square analyses, ANOVAs and t-tests, which illustrate that dizziness and PCS combined are associated with a greater severity of PCS, comorbid depression, and anxiety. Examination of pre-and-post treatment PCS scores showed that both groups improved with treatment.

The treatment protocol used by the mTBI service did remediate PCS in both groups and emerging research indicates that VRT may be effective in reducing PCS and comorbid neuropsychiatric sequelae.³¹⁻³² However, it is difficult to determine to what extent VRT may have affected PCS in the current study, as retrospective analyses cannot provide as robust a research design as that of an RCT with comparative treatment groups and controls. Whilst there were no significant post-treatment differences between those with dizziness and PCS and those with PCS only, it is noteworthy that these groups had comparable post-treatment scores, despite the dizziness and PCS group having much worse symptoms pre-treatment.

In light of the current study's findings, it is suggested that neuro-otological assessments be carried out in all patients with PCS. Forty percent of the current sample were referred to the mTBI service with dizziness complaints and 90% of all mTBI cases were blunt injuries sustained most frequently from falls and playing sports (45% combined). The links between blast-related mTBI and over-pressure trauma to the inner ear are well-established,⁴⁰⁻⁴¹ but secondary blunt injuries from blasts and other blunt mTBI mechanisms can also result in vestibular pathology.⁴² Previous research looking at 5869 UK military personnel deployed to Iraq⁴³ attributed symptoms of post-concussion syndrome to psychiatric disturbance, owing to symptoms being non-specific and there being overlap between presentations. It may not always be practical or possible to provide neuro-otological testing on deployment, however, the possible contribution of vestibular factors to a patient's presentation post-mTBI should be

considered. Indeed, other research from UK military personnel with mTBI has shown an association with dizziness and loss of concentration 7 years post-deployment⁴⁴. It is therefore suggested that there is the potential for inadvertently misattributing vestibular dysfunction and PCS to psychiatric disturbance, thereby overlooking an alternate, more appropriate diagnosis and treatment. Moreover, concussion rehabilitation providers may benefit from engaging the services of VRT specialists sooner so that they can assist with alleviating and treating the myriad of symptoms related to concussion, potentially leading to a faster recovery.

There are some limitations to this retrospective analysis and the use of clinical records. Firstly, we were unable to analyse post-treatment mental health scores, as they were not collected for many of the patients seen through the mTBI service. The analysis also did not account for differences in acute or chronic mTBI presentations which may differ. It was also not possible to account for the potential extent of PTSD and symptom exaggeration influencing PCS and vestibular disorder severity.⁴⁵⁻⁴⁶ Lastly, we were not able to determine the effects of VRT on reducing dizziness symptoms, using retrospective study design. Despite this, there is indication of an intrinsic relationship between the vestibular system, PCS and mental health. Previously, this relationship has shown with co-occurring PTSD to induce devastating long-term functional effects in UK military veterans, where World Health Organisation Disability Assessment Schedule (WHODAS) scores had greater levels of disability than 90% of the general world population.⁵ These conditions have also been linked to a number of pathophysiological and neurodegenerative conditions^{28,47} so should be considered holistically in the context of lifetime adverse health conditions.⁴⁸⁻⁵⁰ Future research should investigate vestibular influences in PCS further and the efficacy of VRT in an RCT that examines both behaviour and biomarkers to establish whether treatment has any long-term effects in remediating PCS sequelae.

In conclusion, dizziness is linked to poorer mental health and greater severity of PCS. However, there are bidirectional links that potentially suggest the vestibular system can both

exacerbate and remediate PCS sequelae. Future research should investigate these relationships holistically in a lifetime context.

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REFERENCES

1. Hawley CA, de Burgh HT, Russell RJ, Mead A. Traumatic brain injury recorded in the UK joint theatre trauma registry among the UK armed forces. *J Head Trauma Rehabil.* 2015;30(1):E47-56. doi:10.1097/HTR.000000000000023.
2. MacGregor AJ, Shaffer RA, Dougherty AL, et al. Prevalence and psychological correlates of traumatic brain injury in Operation Iraqi Freedom. *J Head Trauma Rehabil.* 2010;25(1):1-8. doi:10.1097/HTR.0b013e3181c2993d.

3. Sayer NA. Traumatic brain injury and its neuropsychiatric sequelae in war veterans. *Annu Rev Med.* 2012;63:405-419. doi:10.1146/annurev-med-061610-154046.
4. Kay A, Teasdale G. Head injury in the United Kingdom. *World J Surg.* 2001;25(9):1210-1220. doi:10.1007/s00268-001-0084-6.
5. Denby E, Murphy D, Busuttill W, Sakel M, Wilkinson D. Neuropsychiatric outcomes in UK military veterans with mild traumatic brain injury and vestibular dysfunction. *J Head Trauma Rehabil.* 2020;35(1):57-65. doi:10.1097/HTR.0000000000000468.
6. Dewan MC, Rattani A, Gupta S, et al. Estimating the global incidence of traumatic brain injury. *J Neurosurg,* 2019;130(4):1080-1097. doi:10.3171/2017.10.JNS17352.
7. Lange RT, Brickell TA, Ivins B, Vanderploeg RD, French LM. Variable, not always persistent, postconcussion symptoms after mild TBI in U.S. military service members: a five-year cross sectional outcome study. *J. Neurotrauma.* 2013;30:958–969.
8. Spinos P, Sakellaropoulos G, Georgiopoulos M, et al. Postconcussion syndrome after mild traumatic brain injury in Western Greece. *J Trauma.* 2010;69(4):789-94. doi:10.1097/TA.0b013e3181e3dea67.
9. Sterr A, Herron KA, Hayward C, Montaldi D. Are mild head injuries as mild as we think? Neurobehavioral concomitants of chronic post-concussion syndrome. *BMC Neurol.* 2006;6:7. doi:10.1186/1471-2377-6-7.
10. McDonald SD, Walker WC, Cusack SE, et al. Health symptoms after war zone deployment-related mild traumatic brain injury: contributions of mental disorders and lifetime brain injuries. *Brain Inj.* 2021;35(11):1338-1348. doi:10.1080/02699052.2021.1959058
11. Belanger HG, Vanderploeg RD, Curtiss G, Warden DL. Recent neuroimaging techniques in mild traumatic brain injury. *J Neuropsychiatry Clin Neurosci.* 2007;19(1):5-20. doi:10.1176/jnp.2007.19.1.5.

12. Dash PK, Zhao J, Hergenroeder G, Moore AN. Biomarkers for the diagnosis, prognosis, and evaluation of treatment efficacy for traumatic brain injury. *Neurotherapeutics*. 2010;7(1):100-14. doi:10.1016/j.nurt.2009.10.019.
13. Monsour M, Ebedes D, Borlongan CV. A review of the pathology and treatment of TBI and PTSD. *Exp Neurol*. 2022;351:114009. doi:10.1016/j.expneurol.2022.114009.
14. Thompson SP, McLeod TV. Patient-reported outcomes following vestibular rehabilitation on concussion-induced vertigo: a critically appraised paper. *Int J Athl Ther Train*. 2022;27(5):220-222. doi:10.1123/ijatt.2021-0088.
15. Smith L, Wilkinson D, Bodani M, Bicknell R, Surenthiran SS. Short-term memory impairment in vestibular patients can arise independently of psychiatric impairment, fatigue, and sleeplessness. *J Neuropsychol*. 2019;13(3):417-431. doi:10.1111/jnp.12157.
16. Hoffer ME, Balaban C, Gottshall K, Balough BJ, Maddox MR, Penta JR. Blast exposure: vestibular consequences and associated characteristics. *Otol Neurotol*. 2010;31(2):232-6. doi:10.1097/MAO.0b013e3181c993c3.
17. Pan T, Liao K, Roenigk K, Daly JJ, Walker MF. Static and dynamic postural stability in veterans with combat-related mild traumatic brain injury. *Gait Posture*. 2015;42(4):550-7. doi:10.1016/j.gaitpost.2015.08.012.
18. Banerjee N, Getz SJ, Levin BE. Cognitive-emotional-vestibular triad in mild traumatic brain injury. In: Hoffer ME, Baladan CD, eds. *Neurosensory Disorders in Mild Traumatic Brain Injury*. London: Academic Press; 2019: 183-198.
19. Lempert T, Bronstein A. Management of common central vestibular disorders. *Curr Opin Otolaryngol Head Neck Surg*. 2010 Oct;18(5):436-40. doi:10.1097/MOO.0b013e32833dbd69.

20. Chamelian L, Feinstein A. Outcome after mild to moderate traumatic brain injury: the role of dizziness. *Arch Phys Med Rehabil.* 2004;85(10):1662-6.
doi:10.1016/j.apmr.2004.02.012.
21. Ernst A, Basta D, Seidl RO, Todt I, Scherer H, Clarke A. Management of posttraumatic vertigo. *Otolaryngol Head Neck Surg.* 2005 Apr;132(4):554-8.
doi:10.1016/j.otohns.2004.09.034.
22. Mallinson A, Maire R, Beyaert C, et al. Understanding and managing trauma-induced vestibular deficits. *J Int Adv Otol.* 2021;17(6):559-565. doi:10.5152/iao.2021.21258.
23. Wood NI, Hentig J, Hager M, et al. The non-concordance of self-reported and performance-based measures of vestibular dysfunction in military and civilian populations following TBI. *J Clin Med.* 2022;11(11):2959. doi:10.3390/jcm11112959.
24. Akin FW, Murnane OD. Head injury and blast exposure: vestibular consequences. *Otolaryngol Clin North Am.* 2011;44(2):323-34, viii. doi:10.1016/j.otc.2011.01.005.
25. Bogdanova Y, Verfaellie M. Cognitive sequelae of blast-induced traumatic brain injury: recovery and rehabilitation. *Neuropsychol Rev.* 2012;22(1):4-20.
doi:10.1007/s11065-012-9192-3.
26. Fleminger S. Long-term psychiatric disorders after traumatic brain injury. *Eur J Anaesthesiol Suppl.* 2008;42:123-30. doi:10.1017/S0265021507003250.
27. McAllister TW, Flashman LA, McDonald BC, et al. Dopaminergic challenge with bromocriptine one month after mild traumatic brain injury: altered working memory and BOLD response. *J Neuropsychiatry Clin Neurosci.* 2011;23(3):277-86.
doi:10.1176/jnp.23.3.jnp277.
28. Kaplan GB, Leite-Morris KA, Wang L, et al. Pathophysiological bases of comorbidity: traumatic brain injury and post-traumatic stress disorder. *J Neurotrauma.* 2018;35(2):210-225. doi:10.1089/neu.2016.4953.

29. Hamid MA, Trune DR, Dutia MB. Advances in auditory and vestibular medicine. *Audiol Med.* 2009;7(4):180-188. doi:10.3109/02841860903364076.
30. Haber YO, Chandler HK, Serrador JM. Symptoms associated with vestibular impairment in veterans with posttraumatic stress disorder. *PLoS One.* 2016;11(12):e0168803. doi:10.1371/journal.pone.0168803.
31. Carrick FR, McLellan K, Brock JB, Randall C, Oggero E. Evaluation of the effectiveness of a novel brain and vestibular rehabilitation treatment modality in PTSD patients who have suffered combat-related traumatic brain injuries. *Front Public Health.* 2015;3:15. doi:10.3389/fpubh.2015.00015.
32. Kleffelgaard I, Soberg HL, Bruusgaard KA, Tamber AL, Langhammer B. Vestibular rehabilitation after traumatic brain injury: case series. *Phys Ther.* 2016 Jun;96(6):839-49. doi:10.2522/ptj.20150095.
33. Faul M, Xu L, Wald MM, Coronado V, Dellinger AM. Traumatic brain injury in the United States: national estimates of prevalence and incidence, 2002-2006. *Inj. Prev.* 2010;16(1):A268. doi:10.1136/ip.2010.029215.951.
34. The Management and Rehabilitation of Post-Acute Mild Traumatic Brain Injury Work Group. *VA/DoD clinical practice guideline for the management and rehabilitation of post-acute mild traumatic brain injury.* June, 2021. Accessed July 12th 2023. <https://www.healthquality.va.gov/guidelines/Rehab/mtbi/VADoDmTBICPGFinal508.pdf>.
35. King NS, Crawford S, Wenden FJ, Moss NE, Wade DT. The Rivermead Post Concussion Symptoms Questionnaire: a measure of symptoms commonly experienced after head injury and its reliability. *J Neurol.* 1995;242(9):587-92. doi:10.1007/BF00868811.

36. Spitzer RL, Kroenke K, Williams JB, Löwe B. A brief measure for assessing generalized anxiety disorder: the GAD-7. *Arch Intern Med.* 2006;166(10):1092-7. doi:10.1001/archinte.166.10.1092.
37. Kroenke K, Spitzer RL, Williams JB. The PHQ-9: validity of a brief depression severity measure. *J Gen Intern Med.* 2001;16(9):606-13. doi:10.1046/j.1525-1497.2001.016009606.x.
38. Jacobson GP, Newman CW. The development of the Dizziness Handicap Inventory. *Arch Otolaryngol Head Neck Surg.* 1990;116(4):424-7. doi:10.1001/archotol.1990.01870040046011.
39. Hayes AF. Model templates for PROCESS for SPSS and SAS. In: *Introduction to Mediation, Moderation, and Conditional Process Analysis*. Vol 12. New York, NY: Guildford Press; 2017:2013.
40. Chen Y, Huang W, Constantini S. Concepts and strategies for clinical management of blast-induced traumatic brain injury and posttraumatic stress disorder. *J Neuropsychiatry Clin Neurosci.* 2013;25(2):103-10. doi:10.1176/appi.neuropsych.12030058.
41. Fausti SA, Wilmington DJ, Gallun FJ, Myers PJ, Henry JA. Auditory and vestibular dysfunction associated with blast-related traumatic brain injury. *J Rehabil Res Dev.* 2009;46(6):797-810. doi:10.1682/jrrd.2008.09.0118.
42. Akin FW, Murnane OD, Hall CD, Riska KM. Vestibular consequences of mild traumatic brain injury and blast exposure: a review. *Brain Inj.* 2017;31(9):1188-1194. doi:10.1080/02699052.2017.1288928.
43. Fear NT, Jones E, Groom M, et al. Symptoms of post-concussional syndrome are non-specifically related to mild traumatic brain injury in UK Armed Forces personnel on return from deployment in Iraq: an analysis of self-reported data. *Psychol Med.* 2009;39(8):1379-87. doi:10.1017/S0033291708004595.

44. Rona RJ, Jones M, Fear NT, et al. Mild traumatic brain injury in UK military personnel returning from Afghanistan and Iraq: cohort and cross-sectional analyses. *J Head Trauma Rehabil.* 2012;27(1):33-44. doi:10.1097/HTR.0b013e318212f814.
45. Armistead-Jehle P, Lange BJ, Green P. Comparison of neuropsychological and balance performance validity testing. *Appl Neuropsychol Adult.* 2017;24(2):190-197. doi:10.1080/23279095.2015.1132219.
46. Armistead-Jehle P, Cooper DB, Grills CE, et al. Clinical utility of the mBIAS and NSI validity-10 to detect symptom over-reporting following mild TBI: A multicenter investigation with military service members. *J Clin Exp Neuropsychol.* 2018;40(3):213-223. doi:10.1080/13803395.2017.1329406.
47. Mavroudis I, Kazis D, Chowdhury R, et al. Post-concussion syndrome and chronic traumatic encephalopathy: narrative review on the neuropathology, neuroimaging and fluid biomarkers. *Diagnostics (Basel).* 2022;12(3):740. doi:10.3390/diagnostics12030740.
48. Brown EM, Salat DH, Milberg WP, Fortier CB, McGlinchey RE. Accelerated longitudinal cortical atrophy in OEF/OIF/OND veterans with severe PTSD and the impact of comorbid TBI. *Hum Brain Mapp.* 2022;43(12):3694-3705. doi:10.1002/hbm.25877.
49. Sommer JL, Mota N, Thompson JM, et al. Associations between courses of posttraumatic stress disorder and physical health conditions among Canadian military personnel. *J Anxiety Disord.* 2022;87:102543. doi:10.1016/j.janxdis.2022.102543.
50. Mysliwiec V, McGraw L, Pierce R, Smith P, Trapp B, Roth BJ. Sleep disorders and associated medical comorbidities in active duty military personnel. *Sleep.* 2013;36(2):167-74. doi:10.5665/sleep.2364.