

# Mild Traumatic Brain Injury Classification

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see [Traumatic Brain Injury Classification](#)

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[Concussion](#) and [mild traumatic brain injury](#) (mTBI) are not interchangeable. Concussion may be thought as a subcategory of mTBI on the less severe end of the [brain injury](#) spectrum, though with similar clinical symptoms <sup>1)</sup>.

A major difference between the two is that mTBI may demonstrate abnormal structural imaging (such as [cerebral hemorrhage/contusion](#)) and [concussion](#), by definition, must have normal imaging studies. mTBI is part of an injury severity spectrum primarily based on [GCS](#) score. TBI is evaluated 6 hours after injury and differentiated into mild, moderate and severe

[Concussion](#) is evaluated directly after the insult and based on a clinical diagnosis aided by a multitude of standardized assessment tools. To include concussion under the full spectrum of [traumatic brain injury](#) then it must fall at the low end of mTBI and overlap with the subset of “minimal” injury. Most mTBIs with negative imaging can be considered concussions, but the majority of [sports concussions](#) cannot be classified as mTBI <sup>2) 3)</sup>.

A definition of [Mild Traumatic Brain Injury](#) has been developed by the Head Injury Interdisciplinary Special Interest Group of the American Congress of Rehabilitation Medicine. Within the spectrum of [injury severity](#) in mild TBI there are several [classification](#) systems, primarily used in management of acute mild TBI, that breakdown mild TBI into grades of injury severity. These are based upon the presence or absence of [mental status](#) changes, [amnesia](#), [loss of consciousness](#), anatomical [lesion](#) or [neurological deficit](#) <sup>4)</sup>.

In 1999, a Task Force on Mild Traumatic Brain Injury (MTBI) was set up under the auspices of the European Federation of Neurological Societies. Its aim was to propose an acceptable uniform nomenclature for MTBI and definition of MTBI, and to develop a set of rules to guide initial management with respect to ancillary investigations, hospital admission, observation and follow-up <sup>5)</sup>.

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The objective of a study of Si et al., from the School of Computing, Informatics, and Decision Systems Engineering, Arizona State University, Tempe, AZ, United States of America was to identify “sub-groups” of mild TBI (mTBI) patients based on data available at the time of the initial post-TBI patient evaluation and to determine if the sub-grouping correlates with patient outcomes at 90 and 180 days post-TBI.

Data from patients in the [TRACK-TBI](#) Pilot dataset who had a Glasgow Coma Scale (GCS) score of 13 to 15 at arrival to the Emergency Department and a closed head injury were included. Considering 53 clinical variables that are typically available during the initial evaluation of the patient with mild TBI, sparse hierarchical clustering with cluster quality assessment was used to identify the optimal number of patient sub-groups. Patient sub-groups were then compared for ten outcomes measured at 90 or 180 days post-TBI.

Amongst the 485 patients with mTBI, optimal clustering was based on the inclusion of 12 clinical variables that divided the patients into 5 mild TBI sub-groups. Clinical variables driving the sub-clustering included: gender, employment status, marital status, TBI due to falling, brain CT scan result, systolic blood pressure, diastolic blood pressure, administration of IV fluids in the Emergency Department, alcohol use, tobacco use, history of neurologic disease, and history of psychiatric disease. These 5 mild TBI sub-groups differed in their 90 day and 180 day outcomes within several domains including global outcomes, persistence of TBI-related symptoms, and neuropsychological impairment.

Sub-groups of patients with mTBI can be identified according to clinical variables that are relatively easy to obtain at the time of initial patient evaluation. A patient's sub-group assignment is associated with multidimensional patient outcomes at 90 and 180 days.

The twelve clinical variables that contributed to the clustering structure included: gender, employment status, marital status, injury mechanism, head CT findings, systolic blood pressure, diastolic blood pressure, receiving IV fluids while in the ED, having a history of alcohol use, a history of tobacco use, a history of psychiatric disease, and a history of neurologic disease. Patients in each of the five clusters have different outcomes in regards to global post-TBI outcomes (e.g. GOSE), psychological health (e.g. [BSI](#)), cognition (e.g. WAIS), and post-TBI related symptoms (RPQ). This study helps to identify patient variables that should be further investigated when developing and validating prognostic models for TBI and when identifying more precise sub-categories of mTBI that correlate with patient outcomes. Predictive outcome models consisting of data that are easily and routinely collected during the initial evaluation of patients with mTBI would assist the clinician with determining how aggressively to manage the patient and with providing prognoses to the patients <sup>6)</sup>.

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see [Sport-related concussion](#).

see [Explosive blast mild traumatic brain injury](#).

[Mild Traumatic Brain Injury with traumatic intracranial hematoma](#).

## Pediatric mild traumatic brain injury

[Pediatric mild traumatic brain injury](#).

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2)

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3)

Yuh EL, Hawryluk GW, Manley GT. Imaging concussion: a review. *Neurosurgery*. 2014; 75 Suppl 4: S50–S63

4)

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5)

Vos PE, Battistin L, Birbamer G, Gerstenbrand F, Potapov A, Prevec T, Stepan ChA, Traubner P, Twijnstra A, Vecsei L, von Wild K; European Federation of Neurological Societies. EFNS guideline on mild traumatic brain injury: report of an EFNS task force. *Eur J Neurol*. 2002 May;9(3):207-19. Review. PubMed PMID: 11985628.

6)

Si B, Dumkrieger G, Wu T, Zafonte R, Valadka AB, Okonkwo DO, Manley GT, Wang L, Dodick DW, Schwedt TJ, Li J. Sub-classifying patients with mild traumatic brain injury: A clustering approach based on baseline clinical characteristics and 90-day and 180-day outcomes. *PLoS One*. 2018 Jul 11;13(7):e0198741. doi: 10.1371/journal.pone.0198741. eCollection 2018. PubMed PMID: 29995912.

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