How do we identify the crashing traumatic brain injury patient – the intensivist’s view

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Purpose of review
Over 40% of patients with severe traumatic brain injury (TBI) show clinically significant neurological worsening within the acute admission period. This review addresses the importance of identifying the crashing TBI patient, the difficulties appreciating clinical neurological deterioration in the comatose patient and how neuromonitoring may provide continuous real-time ancillary information to detect physiologic worsening.

Recent findings
The latest editions of the Brain Trauma Foundation’s Guidelines omitted management algorithms for adult patients with severe TBI. Subsequently, three consensus-based management algorithms were published using a Delphi method approach to provide a bridge between the evidence-based guidelines and integration of the individual treatment modalities at the bedside. These consensus statements highlight the serious situation of critical deterioration requiring emergent evaluation and guidance on sedation holds to obtain a neurological examination while balancing the potential risks of inducing a stress response.

Summary
One of the central tenets of neurocritical care is to detect the brain in trouble. The first and most fundamental neurological monitoring tool is the clinical exam. Ancillary neuromonitoring data may provide early physiologic biomarkers to help anticipate, prevent or halt secondary brain injury processes. Future research should seek to understand how data integration and visualization technologies may reduce the cognitive workload to improve timely detection of neurological deterioration.

Keywords
neurocritical care, neurological deterioration, neuromonitoring, secondary brain injury, traumatic brain injury

INTRODUCTION
The detection of secondary brain injury still remains one of the greatest clinical challenges in the management of severe traumatic brain injury (TBI). The development and severity of these secondary injury processes is a major determinant of outcome [1,2]. Consequently, one of the intensivist’s primary responsibilities is to anticipate, prevent and halt secondary brain injury processes during the acute admission period, thereby supporting patients to reach their greatest recovery potential. This is achieved at the bedside through the early detection of clinical neurological deterioration, the maintenance of optimal systemic physiology and the prompt recognition of cerebrovascular pathophysiologic processes using an integrated neuromonitoring approach. The recently published fourth edition of the Brain Trauma Foundation (BTF) guidelines for treating severe TBI in adults consists of high-quality, evidence-based recommendations [3]. However, unlike prior editions [4,5], these guidelines no longer incorporate management algorithms for clinical use. Over the past year, two working groups published three consensus-based management algorithms incorporating expert clinical judgement in areas

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KEY POINTS

- One of the central tenets of neurocritical care is to detect the brain in trouble.
- Up to 44% of patients with severe TBI will develop a clinically relevant neurological worsening during their ICU stay.
- Neuromonitoring is centered on a careful clinical examination, although this can be challenging in comatose and sedated patients.
- Clinicians must balance the decision to hold sedation for clinical examination against the risk of inducing a stress reaction in severe TBI patients, the clinical importance of these stress responses remains to be established.
- A combination of neuromonitoring techniques may provide better insight into brain function than a single monitor used alone.
- In addition to threshold values, trends over time, pressure-time burden and individualized targets are important physiologic concepts to consider when assessing brain function and predicting future neurological deterioration.
- Research into data integration and visualization technologies to optimize high-resolution physiologic data integration may provide new insights into the complex neurophysiologic relationships in critically ill patients with severe TBI and improve earlier detection of neurological deterioration.

wherein current evidence is insufficient. Included in these consensus statements are recommendations to assist in recognizing, evaluating and treating neurological deterioration in patients with severe TBI [6*,7*,8]. This review aims to discuss the recent literature highlighting the importance of detecting the crashing TBI patient, the difficulties identifying neurological deterioration in the comatose patient and the use of neuromonitoring to detect pathophysiologic processes that may act as early biomarkers of neurological deterioration.

NEUROLOGICAL DETERIORATION AFTER SEVERE TRAUMATIC BRAIN INJURY

Despite several decades of basic and clinical research, treatments to improve outcomes after TBI are limited. However, through these trials of neuroprotective therapies, we have gained considerable knowledge about the extent and timing of neurological deteriorations during the acute admission period and associated secondary brain injury processes [9–11]. Secondary brain injury develops over time, from hours to days, with activation of multiple tissue, cellular and molecular pathways (see Fig. 1). Over 40% of patients with severe TBI show significant neurological worsening within the acute admission period [10], warranting immediate medical management and consideration for surgical intervention. The majority of patients deteriorate within 72 h after injury, with a median time of 29 h [11]. In the International Selfotel Trial, the most common neurological deterioration detected was a change in pupillary reactivity (43%), followed by a decrease in the Glasgow Coma Scale (GCS) motor score of more than 1 (25%) [11]. Not surprisingly, patients suffering subsequent neurological deterioration have a significantly higher mortality rate and lower incidence of favourable outcomes than patients with no neurological worsening [9]. Increased intracranial volume accounts for the majority of identified reasons. Interestingly, only a small number of patients deteriorated due to cerebral ischemia and seizures (5 and 7%, respectively) in the International Selfotel Trial, and systemic complications or no definable cause accounted for a quarter of the identified reasons. Interestingly, only a small number of patients deteriorated due to cerebral ischemia and seizures (5 and 7%, respectively) in the International Selfotel Trial, and systemic complications or no definable cause accounted for a quarter of the identified reasons. An understanding of neurological worsening is becoming increasingly important because prompt access to computed tomography (CT) scans within hospitals has resulted in rapid neuroimaging within minutes of admission to hospital before lesions have started to appear or evolve after the primary brain injury [12]. However, parenchymal lesions can expand over hours or days. In a cohort study of 352 patients with brain contusions to investigate the association between clinical and radiological deterioration, the volume of haemorrhage increased in 58% of patients from their first CT at the time of hospital admission to their follow up CTs [13].

DEFINING CLINICALLY SIGNIFICANT NEUROWORSENING

Deterioration of a patient’s clinical status, or neuroworsening, was first defined as a potential intermediate-outcome variable for TBI trials [11]. The International Selfotel Trial defined neuroworsening as the occurrence of one or more of the following objective criteria: a spontaneous decrease in the Glasgow Coma Scale (GCS) motor score of at least 2 points (compared with the previous examination), a new loss of pupillary reactivity, interval development of pupillary asymmetry of at least 2 mm or deterioration in neurological status sufficient to warrant immediate medical or surgical intervention [11]. Neuroworsening was adapted as a clinical variable for the Benchmark Evidence
from South American Trials: Treatment of Intracranial Pressure (BEST:TRIP) trial [14] and further refined by the Consensus REVised Imaging and Clinical Examination (CREVICE) Working Group for their ongoing work investigating the effectiveness of an Imaging and Clinical Examination (ICE) management protocol in resource-limited environments without intracranial pressure (ICP) monitoring [8]. The Seattle Severe Traumatic Brain Injury Consensus Conference (SIBICC) Working Group similarly adapted the definition using the clinical term ‘critical neuroworsening’ to promote recognition that this specific situation is a critical event requiring emergent evaluation and consideration of empiric therapy [6**,7**] (see Table 1).

**Table 1.** Neuroworsening definition adopted by the SIBICC and CREVICE WGs [6**,7**,8]

| Critical Neuroworsening | Sedation Hold Needed | Continuously Monitored |
|-------------------------|----------------------|------------------------|
| Spontaneous decrease in GCS motor score of ≥1 point (compared with the previous examination) | Yes | No |
| New focal motor deficit | Yes | No |
| New decrease/loss of pupillary reactivity | No | No |
| New pupillary asymmetry (≥2 mm) or bilateral mydriasis | No | No |
| Herniation syndrome/Cushing’s triad | No | Yes (ICP, HR, BP, RR) |

BP, blood pressure; CREVICE, Consensus REVised Imaging and Clinical Examination; GCS, Glasgow Coma Scale; HR, heart rate; ICP, intracranial pressure; RR, respiratory rate; SIBICC, Seattle Severe Traumatic Brain Injury Consensus Conference; WG, working group.

**a**The term ‘Critical’ neuroworsening is used specifically by the SIBICC WG to promote its recognition as a critical event and guide expedient evaluation and consideration of empiric therapy.

**b**The modified definition of neuroworsening now includes signs of the herniation syndrome and a lower threshold for GCS motor score (≥1 point).

**c**Pupillary asymmetry quantification of ≥2 mm only utilized in the CREVICE protocol, the SIBICC WG does not quantify the difference in pupillary asymmetry.
DETECTING CLINICAL DETERIORATION WITH SEDATION INTERRUPTION

The first and most fundamental neurological monitoring tool is the repeated clinical examination, even in patients who are comatose or sedated [15]. The minimal requirement for the clinical examination includes an assessment of the level of consciousness, exclusion of new focal neurological deficits and measurement of pupillary size and reactivity to light. The feasibility of using the GCS tool in severe TBI has several potential limitations, confounders such as intoxication, hearing impairment, spinal cord injuries, hypotension, hypoxemia or administration of paralytics compromise assessment [16]. Furthermore, there are obstacles to the assessment of individual components of the GCS such as tracheal intubation precluding a verbal response and ocular trauma impeding eye opening. The motor response remains the main assessable component of the GCS, and fortunately, the prognostic value of the GCS is skewed towards the motor component, as studies have found this single category to be a strong predictor of outcome [17–19]. Sedation interruption, or the neurological wake-up test (NWT), is necessary in deeply sedated patients to evaluate for neurological deterioration, planning of neuroimaging and potential indications for surgical or medical interventions. However, the NWT can induce a stress reaction in severe TBI patients. Prior small single-centre cohort studies have found sedation holds in acutely brain-injured patients can cause transient rises in arterial blood pressure, heart rate, ICP and cerebral perfusion pressure (CPP), an increase in circulating stress hormones and differential results for brain tissue oxygen ($P_{br}O_2$) and cerebral microdialysis (CMD) [20,21,22,23]. The clinical importance of these stress responses remains to be established, as little is known about the impact on patient outcome but should be carefully considered when deciding on the use and frequency of the NWT. The SIBICC Working Group recognized the balance between obtaining the most accurate neurological examination during a sedation hold and the potential hazards of temporarily halting sedation to perform these examinations [7**]. Contraindications to the NWT reported in the literature have included uncontrolled intracranial hypertension, hyperthermia, status epilepticus, barbiturate treatment and acute respiratory distress syndrome (ARDS) [15]. The SIBICC Working Group was unsuccessful at gaining consensus on relative and absolute contraindications for sedation holds and therefore chose to construct decision-support matrices representing the most relevant clinical variables in differing intracranial hypertension scenarios [7**]. The resulting heatmaps reflect the variability among expert clinicians in the perceived safety to perform a sedation hold in ICP-monitored severe TBI patients under differing conditions of pupillary status, GCS motor score, modified Marshall CT classification, duration of ‘controlled’ ICP with ongoing treatment, and degree of tiered therapy required to control any intracranial hypertension. Green, yellow and red indicate ‘safe to proceed’, ‘consider proceeding with caution’ and ‘do not proceed’, respectively, with transitional shades reflecting intermediate trends. Ultimately, it is up to the treating physician to consider the value of performing the NWT, weighing up the risks and benefits. The SIBICC Working Group recommend minimizing risks and enhancing the utility of sedation holds by coordinating the timing for all involved healthcare providers to be present (e.g. Intensivist, Neurosurgeon, ICU nurse) to maximize the safety and interpretation of the NWT [7**]. Overall, sedation holds are not feasible in one-third of patients due to safety concerns and when the NWT is performed, over one-third of these trials are aborted due to critical increases in ICP and impending brain tissue hypoxia [20,24]. Littte is known about the effectiveness of repeated NWTs to detect neurological deterioration and impact on outcomes; however, one small cohort study found that the NWT detected clinical neurological deterioration in one sedation hold out of a total of 54 trials performed [20]. However, given that 29–44% of patients with severe TBI will develop a clinically relevant neurological worsening during their ICU stay, the NWT may help to identify clinically important changes, arguing for repeated neurological examinations (See Table 2 for summary) [10,11,13]. The NWT may have a profound effect on patient management, with aggressive intervention in patients who show signs of progressive brainstem impairment, or reduced duration of ventilation in those recovering favourably.

DETECTING SECONDARY BRAIN INJURY WITH NEUROMONITORING

Due to the concerns of an increased stress response and the energy metabolic challenge to the injured brain, the use of the NWT has been questioned due to the increased access to neuromonitoring. However, the NWT remains the gold standard for clinical monitoring and should always be considered in TBI patients with stable baseline ICP and CPP readings. The overall aims of neuromonitoring are to identify neurophysiologic worsening that may indicate new or ongoing secondary processes, provide clear physiological data to guide and individualize therapy, improve pathophysiological understanding of cerebral disease in critical illness and assist with prognostication [25]. We will focus on the capability of neurophysiologic monitoring to identify neurological deterioration. Neuromonitoring can provide...
ancillary information when assessing which TBI patients can safely undergo the NWT. They also allow a physiologic examination, in place of a clinical examination, when it is either unsafe to perform the NWT in an unstable patient, the severity of the patient’s illness obscures the clinical examination due to level of consciousness or medical interventions, such as an induced coma for intracranial hypertension, prohibit clinical examination. The combined use of multiple brain physiologic monitors, a platform often termed ‘multimodality neuro-monitoring’ [26], can add additional information on brain tissue oxygenation, brain temperature and cerebral metabolism with the aim of providing a continuous, real-time evaluation of the brain’s physiologic state to help prevent, detect and attenuate secondary brain injury [27]. Neuromonitoring devices can be divided into invasive and noninvasive (see Table 3 for an overview). Although the BTF guidelines for treating severe TBI advocate for threshold-based management treating ICP more than 22 mmHg due to the association with increased mortality [3], other emerging important neurophysiologic concepts to consider include the ICP intensity and duration or the ‘pressure-time burden’ [28], ICP trajectory [29] and individualized targets of ICP and CPP [30,31] are important physiologic concepts to consider when assessing brain function and predicting future neurological deterioration. Although clinical studies support the physiologic feasibility and biologic plausibility of monitoring and management based on the information from various cerebral physiologic monitors [25], data supporting this concept from randomized controlled trials are still required. The results of ongoing clinical trials to determine the effectiveness of multimodal neurormonitoring-

Table 2. Large studies reporting timing and features of neurological deterioration

| Study         | Rate of neurological worsening | Time period for neuro-worsening | Neuroworsening criteria | Radiological deterioration |
|---------------|-------------------------------|---------------------------------|-------------------------|---------------------------|
| Iaccarino et al. [13] | 32% (111/352) | Clinical assessment: onset of neurological deterioration during the first 12 hours after trauma | - GCS decreased by >1 point | On follow-up CT scans compared to admission CT |
|               | Clinical improvement in 6%, stable neurological function in 62% | Radiological Assessment: - Injury to initial CT average 120 mins (IQR 63–98 mins) | - New pupillary abnormalities | Patients: 58%: Evolution of hematoma (42% with >30% evolution) |
|               | ‘29% of cohort had severe TBI, other patients had mild and moderate TBI | - 2nd CT average 9 hours after initial scan (IQR 154–312 mins) | | 46%: Increased edema volume |
|               | | - 3rd CT average 38 hours after initial scan (IQR 12–14 hours) | | 30%: Onset/increase in basal cistern effacement |
| Maas et al. [10] | 44% (375/846) | Within first 10 days | Neuroworsening criteria (occurrence ≥ 1 of following): | 28%: Onset /increase of midline shift |
| Morris et al. [11] | 29% (117/409) | Median 29 h [range 3.3–447 h]a | - Spontaneous decrease in GCS motor score ≥2 points (compared with previous exam) | References Morris et al. [11] study for neuroworsening criteria |
|               | | | - New loss of pupillary reactivity | Events: | |
|               | | | - Interval development of pupillary asymmetry of ≥2 mm | 43%: Change in pupillary reactivity | |
|               | | | - Deterioration in neurological status sufficient to warrant immediate medical/surgical intervention | 25%: Decrease ≥ 2 GCS motor score | |
|               | | | Events: | 19%: Pupillary asymmetry >1 mm | |
|               | | | - Changes in ICP | 9%: Changes in ICP | |
|               | | | - Other (decrease GCS ≥ 2, new CT abnormalities, substantial change in systolic BP, systemic deterioration) | 4% - Other (decrease GCS ≥ 2, new CT abnormalities, substantial change in systolic BP, systemic deterioration) |

aData from Juul et al. posthoc analysis of the International Selfotel Trial [9].
BP, blood pressure; CT, computerized tomography; GCS, Glasgow Coma Scale; ICP, intracranial pressure; IQR, interquartile ratio; TBI, traumatic brain injury.
targeted treatment ($P_{btO2}$ and ICP) versus ICP-directed therapy are eagerly awaited [32–34].

DATA VISUALIZATION: INTEGRATING PHYSIOLOGICAL MONITORING TO HELP DETECT CRASHING PATIENTS

As neuromonitoring technology has advanced, the science of data integration and visualization has not kept pace. Tracking, quantifying and displaying dynamic neurophysiologic measures is crucial in the complex care of neurocritically ill patients, and needs to be put into the context of arterial blood pressure, temperature modulation, laboratory results, sedation levels and mechanical ventilator settings, as well as response to other therapeutic interventions. In today’s neurocritical care environment, clinicians are required to assimilate these multiple streams of data in their heads in an attempt to understand the dynamic physiologic interactions between the injured brain and body and detect the brain in trouble. Clinicians are confronted with this high-dimensional data on a daily basis in the neurocritical care unit, with more than 200 data points to review for each patient during the morning ward round [35]. As humans, we find it significantly problematic to remember and simultaneously process data involving more than seven variables [36], and most clinicians are not able to judge the degree of relatedness between more than two variables [37,38]. Along with the introduction of multimodality neuromonitoring into the neurocritical care environment, the ability to acquire biomedical data has outstripped our ability to understand it, all of which greatly contributes to ‘information overload’ that can lead to missed opportunities to detect early physiologic signatures of neurological deterioration and preventable medical errors [27,39]. The fully automated ICU of the future where monitoring technology enables improvements in clinical care has been predicted since the 1970s, but the dream of complete physiologic monitoring captured by a single computer interface has yet to be realized [40,41]. Additional isolated monitoring devices present information on individual smaller displays that may or may not be integrated with the primary patient monitor. In this

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**Table 3. Commonly used neuromonitoring devices**

| Device                                      | Physiological parameter | Global vs. focal physiology | Interpretation/derived indices                                                                 |
|---------------------------------------------|-------------------------|-----------------------------|-------------------------------------------------------------------------------------------------|
| **Invasive neuromonitoring devices**        |                         |                             |                                                                                                |
| ICP monitor (Intraparenchymal/ventricular catheters) | ICP                    | Global                      | Raised intracranial pressure reduces cerebral perfusion                                       |
|                                             |                         |                             | CPP, pressure-reactivity index, intracranial elastance                                         |
| Parenchymal ($P_{btO2}$)                    | Brain tissue partial tension of oxygen | Focal                       | Oxygen diffusion                                                                               |
|                                             |                         |                             | Balance between oxygen supply and demand                                                       |
| Jugular venous oximetry ($S_{jvO2}$)        | Oxygen saturation of jugular haemoglobin | Global                     | Global cerebral oxygenation and extraction                                                     |
|                                             |                         |                             | Cerebral arteriojugular difference in oxygen content                                            |
| Cerebral microdialysis                      | Cerebral metabolism and biomarkers | Focal                       | Aerobic or anaerobic metabolism, brain injury severity and inflammation                        |
| Temperature monitoring (Intraparenchymal probe) | Brain temperature     | Focal                       | Gradient between core and brain temperature                                                    |
| Intraparenchymal thermal diffusion flowmetry | Cerebral blood flow    | Focal                       | Hypoperfusion or hyperperfusion                                                                 |
| **Noninvasive neuromonitoring devices**     |                         |                             |                                                                                                |
| Electroencephalography                      | Cortical electrical activity | Global                     | Seizure activity, abnormal patterns                                                            |
| Optic nerve sheath ultrasonography          | Optic nerve-sheath diameter | Global                     | Elevated value is an indirect marker of raised ICP                                             |
| Quantitative pupillometry                   | ICP                    | Global                      | Low NPI is associated with sustained elevations of ICP                                         |
| Transcranial Doppler                        | Cerebral blood velocity | Focal                       | Indicative of regional cerebral ischemia                                                        |
| Near-infrared spectroscopy                  | Cerebrovascular oxygen saturation | Focal                     | Critical closing pressure, cerebral arterial impedance                                           |

Adapted from Stocchetti et al. [12].
Digital world, neurocritical care clinicians should be able to continually and rapidly evaluate the effect of treatments on the brain and be able to effortlessly track a patient’s vital signs over minutes, hours and days. As fundamental as this may appear for comprehensive neurocritical care, visual plots of multiple waveform data streams that contains core neurophysiologic data are not available at most institutions. However, there is evidence to suggest that even minor improvements in graphical user interfaces such as the presentation of simple line plots of trends or the addition of simple graphical indicators of trend direction could lead to clinically meaningful improvements in diagnostic accuracy and efficiency [42]. Furthermore, a recent systematic review and meta-analysis of 20 studies found that data integration and visualization technologies in critical care were associated with improvements in self-reported performance, mental and temporal demand, and effort compared with paper-based recording systems [43]. However, only 10% of data integration and visualization technology studies evaluated them in clinical settings. Unfortunately, there is a lack of robust evidence on how to integrate and display physiologic monitoring-derived information at the bedside to enhance time to detection of secondary brain injury and improve patient outcomes. The use of a systems design engineering approach to optimize the integration of high-resolution physiologic data will likely provide new insights into the complex neurophysiological relationships in critically ill patients with severe TBI, improve the time to detection of secondary brain injury processes and facilitate the translation of neuromonitoring-driven treatment paradigms.

CONCLUSION AND FUTURE RESEARCH DIRECTIONS

Critically ill patients with severe TBI are at risk of neuroworsening after the initial injury. Studies have shown that patients who subsequently deteriorate during their acute admission period have a much higher mortality and morbidity. The primary role of the intensivist and the critical care interdisciplinary team is the prompt recognition and treatment of any neurological deterioration. As a critical event, this may include empirical medical therapy until further imaging and assessment can be performed. The first monitor will always be the clinical examination, judged by whether it is appropriate to wake up the patient. Further studies are needed to understand whether the clinical information obtained by the NWT justify the risk of inducing a stress response and does this stress response result in subsequent worse longer-term outcomes. Future research should also seek to understand if the use of integrative neuromonitoring facilitates prompt recognition of earlier pathophysiology deterioration and improves outcome, and whether novel data visualization techniques facilitate a better understanding of complex physiological relationships and improved care at the bedside.

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Papers of particular interest, published within the annual period of review, have been highlighted as:

* of special interest
** of outstanding interest

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