Non-invasive estimation of cerebral perfusion pressure using transcranial Doppler ultrasonography in children with severe traumatic brain injury

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Abstract
Objective To identify if cerebral perfusion pressure (CPP) can be non-invasively estimated by either of two methods calculated using transcranial Doppler ultrasound (TCD) parameters.
Design Retrospective review of previously prospectively gathered data.
Setting Pediatric intensive care unit in a tertiary care referral hospital.
Patients Twenty-three children with severe traumatic brain injury (TBI) and invasive intracranial pressure (ICP) monitoring in place.
Interventions TCD evaluation of the middle cerebral arteries was performed daily. CPP at the time of the TCD examination was recorded. For method 1, estimated cerebral perfusion pressure (CPPe) was calculated as: CPPe = MAP × (diastolic flow (Vd)/mean flow (Vm)) + 14. For method 2, critical closing pressure (CrCP) was identified as the intercept point on the x-axis of the linear regression line of blood pressure and flow velocity parameters. CrCP/CPPe was then calculated as MAP-CrCP.
Measurements and main results One hundred eight paired measurements were available. Using patient averaged data, correlation between CPP and CPPe was significant (r = 0.78, p < 0.001). However, on Bland-Altman plots, bias was 3.7 mmHg with 95% limits of agreement of −17 to +25 for CPPe. Using patient averaged data, correlation between CPP and CrCP/CPPe was significant (r = 0.59, p < 0.001), but again bias was high at 11 mmHg with wide 95% limits of agreement of −15 to +38 mmHg.
Conclusions CPPe and CrCP/CPPe do not have clinical value to estimate the absolute CPP in pediatric patients with TBI.

Keywords Cerebral perfusion pressure • Transcranial Doppler ultrasound • Ultrasound • Head injury • Non-invasive monitoring • Traumatic brain injury

Cerebral perfusion pressure (CPP) is the driving pressure gradient which produces blood flow in the cerebral circulation. Thus, CPP is calculated as the difference between the mean cerebral arterial blood pressure and the effective downstream pressure or the critical closing pressure (CrCP) (CrCP = tissue pressure + cerebral vascular tone + mean cerebral venous pressure). As it is difficult to measure these parameters necessary to calculate CrCP, the current gold standard to determine CPP clinically is to subtract the invasively monitored intracranial pressure (ICP) from the mean systemic arterial pressure (MAP) (CPP = MAP − ICP).

In critical illness, systemic hypotension or intracranial hypertension may lead to a reduction in CPP and result in secondary neurologic insult. Additionally, when cerebral autoregulation is impaired or absent, as occurs in many disease states, the risk of these secondary insults increases as cerebral blood flow becomes linearly correlated with CPP [1–7]. As such,
monitoring and optimizing CPP as a targeted therapy has shown promise to improve outcomes in children with critical illness [8–13]. Management guidelines for children with severe traumatic brain injury (TBI) recommend clinicians consider, as an option, maintaining minimum age-specific CPP goals [12]. In 317 children with TBI, a significant increase in mortality risk was noted when CPP fell below the desired level (relative risk of death (RR) = 8.1 (95% CI 3.58, 18.31)) [8]. In childhood bacterial meningitis, CPP-directed therapy has been shown to reduce overall mortality by 18% (RR = 2.1; 95% CI, 1.09–4.04; p = 0.02) and neurodisability by 33% (RR = 0.47; 95% CI, 0.27–0.83; p = 0.004) when compared with ICP directed therapy alone [10].

However, placement of an ICP monitor that allows for the calculation of CPP does not always occur in the clinical care of these children. Neurosurgical expertise necessary to place invasive monitors is often not available in rural or in resource-limited settings. Additionally, in some centers, there remains a lack of confidence that ICP/CPP-directed therapy benefits patients and children are managed with alternative therapeutic approaches [14–16]. Furthermore, while the risk is minimal, it is an invasive surgical technique occasionally associated with complications including infection and hemorrhage [17]. In a cohort of 4667 children with severe TBI, only 55% of patients had an ICP monitor placed [18]. Younger children may be even less likely to receive invasive ICP monitors. In another study of 238 children under age 24 months with severe TBI, only 17% had a monitor placed [19]. Outside the management of TBI, use of an ICP monitor in children may be even rarer. Odetola et al. evaluated > 1000 children requiring mechanical ventilation during treatment for severe bacterial meningitis, and found placement of an ICP monitor occurred in only 7% of cases [20]. Other examples where CPP monitoring may be helpful but in whom invasive monitoring is not routinely performed or is contraindicated include children with coagulopathy in the setting of sepsis or liver failure, or while undergoing support with extracorporeal membrane oxygenation [21–24].

Identifying a non-invasive, simple means by which CPP can accurately and reliably be quantified and monitored in children may be a helpful adjunct to care when treating clinicians cannot or do not place an invasive monitor but have an ongoing desire to evaluate cerebral perfusion. Transcranial Doppler ultrasound (TCD) is a portable, repeatable, non-invasive means that measures cerebral flow velocities (CBFVs) in the cerebral vasculature. In adult patients, TCD-derived parameters have successfully been used to calculate an estimated CPP by several different mathematical approaches [25–29]. Two that have been described require only systemic blood pressure monitoring and basic non-continuous TCD data [27, 30–32]. One method described by Czosnyka et al. calculates an estimated CPP (CPPe) as MAP × (diastolic flow (Vd)/mean flow (Vm)) + 14 [27]. In the second method, cerebral CrCP can be determined by plotting the dynamic pressure-flow relationship between systemic blood pressures and TCD flow velocities and determining the x-axis intercept of the linear regression line [30–32]. Estimated CPP by this method (CrCP/CPPe) is then calculated as MAP − CrCP [30–32]. There is a paucity of literature evaluating the use of either of these methods to non-invasively estimate CPP in pediatric patients. We therefore performed the current study to test the hypothesis that CPPe and CrCP/CPPe derived according to these methods would have good agreement with invasively monitored CPP in children.

Materials and methods

Study population

We performed a retrospective review of data previously gathered from 2011 to 2015 for a prospective, observational study in a tertiary care pediatric intensive care unit. This study was approved by the Institution Review Board. Children 1 day to 17 years of age admitted with a diagnosis of severe TBI (post-resuscitation Glasgow Coma Scale (GCS) score ≤ 8) managed with an invasive ICP monitor (intraparenchymal monitor, Camino Integra Neurosciences, Plainsboro, NJ, USA) and radial arterial line were included. Patients with non-traumatic etiologies of admission were excluded. Demographic data including age, gender, GCS score, and mechanism of injury were recorded for all participants.

General management protocol

Patients were treated following the Society of Critical Care Medicine Guidelines [12]. Patients with surgical lesions underwent resection of the lesion and were left with a primary decompressive craniotomy. Secondary decompressive craniotomy for refractory intracranial hypertension was not performed in our institution during the time period the study was undertaken. All patients received sedation and anxiolysis with infusions of fentanyl and versed. Elevation in ICP (≥ 20 mmHg) despite adequate sedation was treated with osmolar therapy followed by neuromuscular blockade. CPP goals (CPP > 40 mmHg for children < 1 year of age, 50–60 mmHg for children < 12 years of age, and > 60 mmHg for adolescents) were set by the treatment team and were maintained using fluid boluses to a central venous pressure > 10 cmH2O followed by a vasoactive infusion.

Transcranial Doppler ultrasonography

TCD was performed at the participant’s bedside by one of two sonographers using a 2-MHz pulsed probe and commercially available ultrasonography unit (Sonara Digital TCD,
CareFusion, Middleton, WI). The quality of the data obtained by TCD is highly influenced by operator-dependent factors such as skill and experience. Prior to the study beginning, sonographers were tested on standardized patients until a coefficient of variation < 10% for each study measurement was demonstrated. Middle cerebral arteries (MCAs) were insonated at 2-mm intervals using methods previously described, and CBFVs including the Vs, Vd, and Vm were recorded [25, 26]. All participants underwent the initial TCD within 24 h of injury and then daily thereafter through death or hospital day 8, whichever came first. Continuous TCD monitoring was not available at the time of this study.

**Data capture and calculations**

Arterial and intracranial pressure transducers were calibrated at the level of the skull. Outputs from the pressure monitors and the TCD unit (maximal frequency envelope) were connected to an analog-to-digital converter that was fitted into a laptop computer. Data were sampled (sampling frequency 50 Hz), digitized (12 bits), processed, and stored on the computer using software designed in-house for this purpose. Time-averaged (mean) values of pressures were calculated using time integration of waveforms for 5-s intervals. Time-averaged Vs, Vd, and Vm were calculated after spectral filtration to reduce the influence of noise and averaged within 5-s periods. Digital Fourier analysis was used to correlate corresponding values of arterial blood pressure and CBFV. CPPe was then calculated as CPPe = MAP × Vd/Vm + 14 [27]. CrCP was determined as the intercept point of a regression line between arterial systolic and diastolic pressures plotted along the x-axis and the systolic (Vs) and Vd flow velocities plotted along the y-axis [28–30]. CrCP/CPPe was then calculated as CrCP/CPPe = MAP − CrCP.

**Statistical analysis**

Descriptive values were expressed as frequencies for dichotomous variables and as mean ± standard deviation (SD) or median (interquartile range) for continuous variables. To assess the performance of the proposed methods, the correlation between CPP and CPPe and CrCP/CPPe was evaluated using the Spearman correlation coefficient (r, with the level of significance set at 0.05). In addition, a generalized estimating equation was used to account for inter- and intraindividual differences between studies using original observations. Bland-Altman plots were constructed to study agreement between simultaneously invasively measured CPP and TCD derived CPPe (both CPPe and CrCP/CPPe). Two-by-two contingency tables were performed to determine sensitivity, specificity, positive predictive value, and negative predictive value. Statistical significance was assumed with a p ≤ 0.05.

Using patient-averaged data, correlation between CPP and CPPe was significant (r = 0.78, p ≤ 0.001) (Fig. 1). However, when evaluating agreement between CPP and CPPe with a Bland-Altman plot, the bias or average discrepancy for all measurements was 3.7 mmHg with 95% limits of agreement of −17 and +25 mmHg (Fig. 1). When evaluating agreement over time, average discrepancy was best on post-injury day 1 and worsened over time (Table 2). The wide limits of agreement were relatively unchanged by day (Table 2). Fifty-three percent of all CPPe measurements were ≥10 mmHg below or above the invasively calculated CPP.

**CPP and CrCP/CPPe**

Using patient-averaged data, correlation between CPP and CrCP/CPPe was significant (r = 0.59, p <0.001) (Fig. 2). However, when evaluating the agreement between CPP and

**Table 1** Clinical and laboratory data of patients (n = 23)

| Characteristic                        | Result     |
|---------------------------------------|------------|
| Age in months (mean ± SD)             | 96 ± 60    |
| Male (%)                              | 18 (78%)   |
| Mechanism of injury (%)               |            |
| Fall                                  | 2 (9%)     |
| Motor vehicle accident                | 10 (43%)   |
| Pedestrian vs auto                    | 6 (26%)    |
| Abusive head trauma                   | 3 (13%)    |
| Other                                 | 2 (9%)     |
| Glasgow Coma Score (median/IQR)       | 5 (3, 7)   |
| PaCO2 (mmHg) at time of TCD           | 39 ± 6     |
| ICP (mmHg, median/IQR) at time of TCD | 10 (8, 15) |
| CPP (mmHg, median/IQR) at time of TCD | 68 (60, 79) |
| Decompressive craniotomy (%)          | 8 (35%)    |
| In-hospital mortality (%)             | 3 (13%)    |

SD standard deviation, vs versus, PaCO2 partial pressure carbon dioxide, ICP intracranial pressure, CPP cerebral perfusion pressure, TCD transcranial Doppler ultrasound, mmHg millimeters mercury.
CrCP/CPPe with a Bland-Altman plot, the bias was 11 mmHg and 95% limits of agreement were −15, +38 mmHg (Fig. 2). Average discrepancy was high, and limits of agreement were wide on all days (Table 3). Forty-three percent of CrCP/CPPe measurements were ≥10 mmHg below or above the invasively calculated CPP.

Estimated CPP as a screening tool for low CPP

There were no TCD examinations performed during episodes of CPP < 40 mmHg available for analysis. There were 14 episodes of CPP < 50 mmHg when the age specific goal was > 50 mmHg (mean CPP 44 ± 2 mmHg) and 22 episodes of CPP < 60 mmHg when the age specific goal was > 60 mmHg (mean CPP 51 ± 3 mmHg). Sensitivity, specificity, positive predictive value, and negative predictive value of both CPPe and CrCP/CPPe to detect reduced CPP below treatment goal values are in Table 4.

Confounders

Given the potential changes to intracranial compliance following surgical intervention, the agreement between CPP and CPPe as well as CPP and CrCP/CPPe was assessed in children who underwent a primary decompressive craniectomy (n = 8) separately than in those who did not (n = 15). The Bland-Altman plot testing agreement between CPP and CPPe in...
children with decompressive craniotomy found bias was 5 mmHg with 95% limits of agreement ranging from −56 to 45 mmHg. In children without decompressive craniotomy, bias was 2 mmHg and 95% limits of agreement were −28 to 32 mmHg. The Bland-Altman plot evaluating agreement between CPP and CrCP/CPPe in children with and without decompressive craniotomy revealed a bias of 15 mmHg with 95% limits of agreement of −22 to 49 mmHg and bias of 13 mmHg with 95% limits of agreement of −12 to 45 mmHg, respectively.

Bland-Altman plots were also performed to evaluate the effect of partial pressure of carbon dioxide (PaCO₂) on the agreement between CPP and CrCP/CPPe. At PaCO₂ values of 30–35 mmHg (n = 28), bias was −0.15 mmHg and 95% limits of agreement were −27 to 27 mmHg. At PaCO₂ values of 35–40 mmHg (n = 37), bias was 2.5 mmHg with 95% limits of agreement −18 to 23 mmHg. At PaCO₂ values of 40–45 mmHg (n = 30), bias was 5.9 mmHg with limits of agreement at −17 and 29 mmHg. At PaCO₂ > 45 mmHg (n = 13), bias was 6.1 mmHg with 95% limits of agreement at −17 and 26 mmHg. Values for agreement between CPP and CrCP/CPPe were similar at different PaCO₂ ranges in terms of limits of agreement (data not shown).

Minor observations

Three patients had repeated, real-time TCD recordings before, during, and after physiologic derangements that allowed for the calculation of serial CrCPs. In one patient, ICP spontaneously increased from 12 to 24 mmHg. MAP changed minimally from 84 to 86 mmHg. Pre-ICP spike, CrCP was calculated as 19 mmHg and then increased to 39 mmHg during the ICP spike. CrCP/CPPe was thus 65 mmHg at baseline and decreased to 47 mmHg during the ICP increase. Five milliliters (ml)/kg of 3% hypertonic saline were given and ICP decreased to 13 mmHg and MAP increased to 90 mmHg. CrCP/CPPe thus increased to 71.2 mmHg [33, 34]. During the ICP spike, end-tidal CO₂ monitoring did not significantly change and there was no clear increase in cerebral metabolism (patient was undergoing sedation and neuromuscular blockade and electroencephalogram did not reveal seizures). Another patient had similar pathologic derangements with similar alterations to calculated CrCP and CrCP/CPPe during ICP spikes. A third patient experienced refractory intracranial hypertension with an ICP of 45 mmHg. The patient was given 1 g/kg of mannitol, 5 ml/kg of hypertonic saline, and mechanical ventilator rate was increased to result in an end tidal CO₂ reduction from 35 to 30 mmHg. Despite these interventions, ICP increased further to 52 mmHg. CrCP was 60 mmHg when the ICP was 45 mmHg and increased further to 72.9 mmHg as ICP increased (Fig. 4). During this time, the MAP did not significantly change (112 to 113 mmHg) so the CrCP/CPPe decreased from 52 to 40 mmHg.

Discussion

In critically ill children with a variety of primary diagnoses, maintenance of an age-appropriate CPP is suggested as a treatment option to ensure necessary substrate delivery and avoid secondary neurologic insult. However, direct ICP monitoring that allows for calculation of CPP is not always feasible given lack of resources or clinical contraindications. Thus, finding a non-invasive alternative method by which to assess and monitor CPP is desirable and may aid in the management of these children.

Mathematical models using combinations of systemic blood pressure measurements and TCD-derived flow velocities have been suggested to have clinical utility to non-

| Day | Bias | −1.96 SD | +1.96 SD |
|-----|------|----------|----------|
| 1   | 13.4 | −16      | 43       |
| 2   | 14.2 | −19      | 48       |
| 3   | 16.7 | −12      | 44       |
| 4   | 15.9 | −16      | 42       |
| 5   | 16.9 | −14      | 39       |
| 6   | 16   | −17      | 41       |
| 7   | 17.2 | −18      | 42       |
| 8   | 16.6 | −16      | 43       |

mmHg millimeters mercury, SD standard deviation

| CPPe < 50 mmHg | CrCP/CPPe < 50 mmHg | CPPe < 60 mmHg | CrCP/CPPe < 60 mmHg |
|----------------|---------------------|----------------|---------------------|
| Sensitivity    | 100                 | 66             | 73                  | 67                  |
| Specificity    | 91                  | 53             | 78                  | 63                  |
| Positive predictive value | 27               | 8              | 60                  | 48                  |
| Negative predictive value | 100              | 96             | 90                  | 100                 |

CPPe estimated cerebral perfusion pressure, CrCP/CPPe critical closing pressure derived cerebral perfusion pressure
invasively estimate CPP in adult patients [27–29, 32–39]. Given the paucity of literature on this topic in pediatric patients, we performed the current study to evaluate the utility of two methods to non-invasively estimate CPP in children with severe TBI. Our findings include the following: (1) CPPe had adequate average discrepancy from the invasively measured CPP, but had wide limits of agreement at −17 to +25 mmHg; (2) CrCP/CPPe had high average discrepancy from the invasively measured CPP and had wide limits of agreement at −15 to 38 mmHg; (3) the ability of either method to predict CPP < 50 mmHg or < 60 mmHg was insufficient for clinical use; and (4) in a very small number of children, changes to calculated CrCP and CrCP/CPPe over time appropriately reflected underlying pathophysiologic alterations to ICP and CPP and the response of these values to interventions.

Using the first method, we evaluated, Czosnyka et al. determined the correlation between CPP and CPPe was $r = 0.73$, $p < .0001$ with an overall error < 15 mmHg in 84% of the examinations [27]. Schmidt et al. found, using the same formula, that the absolute difference between CPP and CPPe was < 10 mmHg in 89% of measurements with a 95% confidence range for prediction of the actual CPP no wider than ±12 mmHg [28]. Using the second method we evaluated, CrCP/CPPe had good correlation with CPP in 70 adult patients ($r = 0.92$) [31]. These authors promoted the use of both of these methods to non-invasively estimate CPP in adult patients based on a good correlation between the invasively measured and estimated CPP. In fact, correlation of CPP and CPPe from our cohort was also reasonably good ($r = 0.78$). However, since correlation studies the relationship between one variable and another but not the differences between them, it may not be the most appropriate method to assess comparability between a gold standard and new method.

A Bland-Altman plot compares the agreement of two measurement techniques by plotting the difference of two paired measurements against the mean of two measurements. Results then quantify the mean difference between the two methods and give 95% limits of agreement. In adult studies of TBI, the agreement between invasively and non-invasively calculated CPP on Bland-Altman plots by either of these methods is generally wide [31, 32, 35]. These results likely explain why, despite the technique and technology now having been available for more than 20 years, the approach has not routinely been adopted into clinical practice for adults.

In a recently published paper using continuous TCD recordings to non-invasively estimate CPP in children who had suffered TBI, CPPe overestimated CPP by 19.61 mmHg with wide 95% CI of ±40 mmHg on Bland-Altman analysis [36]. Our results are similar to this study in that we also identified wide limits of agreement for both the CPPe and CrCP/CPPe methods (>−15 to >+25). Thus, the clinical utility of either method to non-invasively estimate the absolute value of CPP at a single time point in children with TBI also appears to be limited.

Some work has previously been done suggesting that TCD parameters may have a role in non-invasively estimating CPP < 50 mmHg in children with severe TBI [40]. Figaji et al. reported that the TCD-derived PI was 0.95 ± 0.17 when CPP was ≤50 mmHg and 0.78 ± 0.20 when CPP was ≥50 mmHg. However, sensitivity and specificity of the PI to determine a
CPP below 50 mmHg was not done due to an inadequate number of paired measurements when CPP was low. In our study, in a limited number of episodes \( (n = 14) \), CPPe was noted to have a sensitivity of 100%, specificity of 91%, and a positive predictive value of 27% to predict CPP < 50 mmHg. The low positive predictive value we report is likely due to the relatively low prevalence of cerebral hypoperfusion < 50 mmHg in this cohort of children. Larger studies can be considered to evaluate if these values of sensitivity and specificity for any of these methods hold with increased measured episodes of cerebral hypoperfusion < 50 mmHg. However, the significant reduction in sensitivity and specificity of this method in a larger number \( (n = 22) \) of subjects at a higher target CPP (< 60 mmHg) seen in our study suggests that they likely will not. Identifying a non-invasive means by which to determine cerebral hypoperfusion is thus still necessary.

The study revealed modestly improved agreement between the estimated and invasively measured CPP in children who had not undergone decompressive craniotomy compared with those that had. As ICP rises and reaches or exceeds the critical closing pressure at the arteriolar level, other contributors to the calculated CrCP become less important and the discrepancy between the estimated and actual CPP decreases [41]. Following decompressive craniotomy, ICP in general is assumed to not reach critical levels. Sample sizes did not allow for matched comparisons at various ICP values between children with and without DC, but a difference in ICP/higher ICP may have contributed to the slightly improved agreement in patients who had not undergone DC. However, wide limits of agreement in this group of children would still prohibit its use clinically. Additionally, bias between the methods was slightly better when the PaCO\(_2\) values were low (30–35 mmHg) and progressively worsened as PaCO\(_2\) values rose (> 45 mmHg). Autoregulation is known to be most effective and CVR highest at lower PaCO\(_2\) values [42, 43]. The overall effect of these factors on measured CBFVs may account for the modest improvement in agreement at relative hypocapnea.

Another important clinical need is the ability to monitor non-invasively, not just at a single point but over time, the actual cerebral perfusion of these critically ill children. The components that contribute to the calculation of the CrCP, and thus, the CrCP/ CPPe include not just the tissue pressure (ICP) but also the cerebrovascular resistance/vasomotor tone (CVR), and the downstream venous pressure [30, 31]. Thus, the calculation of the CrCP/CPPe reflects the net effect of local and systemic physiologic or pathologic alterations on the cerebral circulation. If trended over time, it may be useful to assist in determining an individual’s overall response to progressive or improving pathologic states and the effects of therapeutic strategies on the effective cerebral perfusion pressure. In two small experimental animal studies, this has been done successfully. Varsos et al. calculated CrCP during 38 episodes of ICP plateau waves induced by lumbar infusion. ICP increased during infusion on average by 24 mmHg and a concomitant increase in CrCP of 27% from baseline was identified (mean CrCP 51.9 ± 8.76 at baseline ICP; mean CrCP 63.31 ± 10.83 at the top of the plateau waves) [34]. The same author, in a different study, determined that alterations to the calculated baseline CrCP and CrCP/CPPe accurately reflected the overall net effect of changes to ICP, mean arterial pressure, and ventilation in rabbits [44]. Three patients in our cohort had serial TCD examinations around the time of significant pathophysiologic changes and treatment interventions that allowed for the repeated calculation of CrCP and thus CrCP/CPPe. In all 3 children, CrCP and CrCP/CPPe trended in the expected direction based on the measurement of simultaneously captured invasive parameters. This is a very limited sample, however, so the importance of these results must not be overstated. Future studies in larger numbers of children should determine the utility of CrCP calculated non-invasively in this way to monitor for changing physiology and as a means to measure the response to therapeutic interventions in children with acute, severe, neurologic illness.

**Limitations**

This study involved a relatively small cohort of patients who experienced a limited range of CPP variations. Due to the limited number of participants, we were unable to determine if there were certain patient characteristics that determined good versus poor agreement between the presented methods and the invasively measured CPP. Future studies could involve larger groups of children and attempt to capture agreement data with different subsets of patients and determine if CPPe or CrCP/ CPPe can be used to non-invasively predict CPP in some children. Furthermore, this study did not evaluate the techniques to non-invasively estimate CPP in children with disease processes outside TBI. Other disease processes that have different pathophysiologic mechanisms and more diffuse versus heterogeneous neurologic injury may have different levels of agreement.

If any of these studies were to find closer agreement between estimated CPP and invasively measured CPP, other considerations would need to be taken into account before widespread clinical use of the technique began. One limitation of TCD use in general is the considerable variation that can be seen between different operators. It would therefore be imperative that all operators at a single institution would have to undergo regular evaluation to ensure minimal variation in technique and results of their TCD examinations. This would help to ensure accurate and interpretable results when repeated examinations are done on a single patient over time by multiple operators. Furthermore, as CPP is highly dynamic and can fluctuate widely within minutes, improved technology that allows for continuous reliable TCD measurement and subsequent real-time non-invasive CPP calculation in children would be required in order to use the technique as anything other than a one-time screening tool.
Conclusions

CPPe and CrCP/CPPe do not have clinical values in the non-invasive estimation of the absolute CPP measured invasively or to detect cerebral hypoperfusion below desired thresholds of 50–60 mmHg in children with severe TBI.

Compliance with ethical standards

This study was approved by the Institution Review Board.

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