Therapeutic hypothermia in patients with coagulopathy following severe traumatic brain injury

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Abstract

Background: Coagulopathy in traumatic brain injury (TBI) has been associated with poor neurological outcomes and higher in-hospital mortality. In general principle of trauma management, hypothermia should be prevented as it directly worsens coagulopathy. Therefore, we examined the safety of mild therapeutic hypothermia (MTH) in patients with coagulopathy following severe TBI.

Methods: We re-evaluated the brain hypothermia (B-HYPO) study data based on coagulopathy and compared the Glasgow Outcome Scale scores and survival rates at 6 months using per protocol analyses. Coagulopathy was defined as an activated partial thromboplastin time (APTT) > 60 s and/or fibrin/fibrinogen degradation product levels (FDP) > 90 μg/mL on admission. Baseline characteristics, coagulation parameters, and outcomes were compared between the control and MTH groups with or without coagulopathy.

Results: In patients with coagulopathy, 12 patients were allocated to the control group (35.5–37.0 °C) and 20 patients to the MTH group (32–34 °C). In patients without coagulopathy, 28 were allocated to the control group and 59 patients were allocated to the MTH group. In patients with coagulopathy, favorable neurological outcomes and survival rates were comparable between the control and MTH groups (33.3% vs. 35.0%, P = 1.00; 50.0% vs. 60.0%, P = 0.72) with no difference in complication rates. On admission, no significant differences in APTT or FDP levels were observed between the two groups; however, APTT was significantly prolonged in the MTH group compared to the control group on day 3.

Discussion: Based on our study, MTH did not seem to negatively affect the outcomes in patients with coagulopathy following severe TBI on admission; therefore, the present study indicates that MTH may be applicable even in patients with severe TBI and coagulopathy.

Conclusions: Our study suggests that in comparison to control, MTH does not worsen the outcome of patients with coagulopathy following severe TBI.

Trial registration: UMIN-CTR, No. C000000231, Registered 13 September 2005.

Keywords: Coagulopathy, Therapeutic hypothermia, Traumatic brain injury, Targeted temperature management, Fibrinogen degradation products

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Background

Coagulopathy in traumatic brain injury (TBI) has been associated with poor neurological outcomes and higher in-hospital mortality [1–5]. However, in previous studies, the reported incidence of coagulopathy in isolated TBI patients has varied from 7% to 86% due to the use of differing definitions of coagulopathy [1, 6, 7].

Recently, a revised lethal triad has been proposed, with coagulopathy (fibrin/fibrinogen degradation products (FDP) levels >90 μg/mL) posited to have a central role [8]. As a general principle of trauma management, hypothermia should be prevented as it directly worsens coagulopathy. In vitro studies have demonstrated that hypothermia below 33 °C can cause coagulation dysfunction; however, the risk of bleeding associated with mild therapeutic hypothermia (MTH) is considered to be relatively small [9, 10].

Many randomized clinical trials (RCT) have been conducted to investigate the effectiveness of MTH for TBI, but they could not demonstrate more favourable outcomes than those obtained by normothermia (at 37 °C) [11–13]. However, the latest guidelines from an expert panel suggested considering TTM at 34–35 °C in order to lower ICP in TBI patients with refractory intracranial hypertension despite medical treatments [14]. Furthermore, trials examining efficacy and safety in patients with coagulopathy following TBI are yet to be conducted [11–13, 15, 16]. Therefore, we examined the hypothesis that MTH is harmful in patients with coagulopathy following severe TBI. The purpose of the present study was to examine the effect of coagulopathy on the safety of MTH compared to control in patients with severe TBI.

Methods

B-HYPO study

The B-HYPO study was conducted as a prospective, multicenter RCT between December 2002 and September 2008. The protocol was approved by the Institutional Review Board of each participating hospital, and the trial was registered at the University Hospital Medical Information Network site (UMIN-CTR, No. C000000231, Registered 13 September 2005) in Japan and at the National Institutes of Health site (Clinical Trials. Gov, Identifier NCT00134472, Registered 23 August 2005) in the United States of America. In brief, inclusion criteria were as follows: age 15–69 years for both sexes and a Glasgow Coma Scale (GCS) score of 4–8. Written informed consent was obtained from legally authorized representatives of patients prior to inclusion. If informed consent could not be obtained within 2 h of admission, the consent policy was waived.

Targeted temperature management (TTM)

Treatments were performed as described in our original paper [15]. In brief, cooling was initiated within 2 h of the onset of TBI. The goal in each group was to achieve the targeted temperature within 6 h of the onset of TBI and to maintain this temperature for at least 72 h, predominantly using surface cooling blankets. After 72 h, the temperature was maintained at <38 °C until 7 days after the onset of TBI.

Definition of coagulopathy

In previous literature, definitions of coagulopathy using the activated partial thromboplastin time (APTT) have varied from 32 s to 60 s [1, 17]. In the present study, we adopted the most severe APTT criteria (60 s). Accordingly, patients with an APTT >60 s and/or an FDP level > 90 μg/mL on admission were allocated to the coagulopathy group [18–20].

Patients

In the original paper, 150 patients were randomly assigned (1:2 allocation ratio) to either the control group (35.5–37.0 °C) or the MTH group (32.0–34.0 °C), and analyzed by intention to treat analyses [15]. Per-protocol analyses were performed in 135 patients (control, 47 patients and MTH, 88 patients) [21]. In the present post hoc study, we re- evaluated these data (n = 135) based on initial coagulation markers, APTT, and/or FDP levels. Sixteen patients (control, 7 patients and MTH, 9 patients) were excluded as either APTT or FDP values were unavailable (Fig. 1). Patients were classified as either coagulopathy (n = 32, 26.9%) or non-coagulopathy (n = 87, 73.1%). In patients with coagulopathy, 12 patients were allocated to the control group and 20 patients to the MTH group. In patients without coagulopathy, 28 were allocated to the control group and 59 patients were allocated to the MTH group.

Data collection and study outcomes

Data on the following parameters were collected: age, gender, systolic blood pressure, heart rate, GCS, unresponsive pupil or pupils, platelet counts, APTT, fibrinogen, FDP, Traumatic Coma Data Bank classification, Injury Severity Score (ISS), Abbreviated ISS (AIS) for the head, AIS score ≥ 4 for other organs on admission, complication rate during TTM, surgical intervention for TBI during administration, and favorable neurological outcomes and survival rates at 6 months following TBI. APTT and FDP levels in the acute phase (time to admission to day 3) were compared between the control and MTH groups with or without coagulopathy, respectively. Platelet counts in the acute phase (time to admission to day 3) were also compared between the control and MTH groups with coagulopathy.

The primary outcomes were favorable neurological outcomes, survival rates, and complication rates between
Comparison of baseline characteristics, neurological outcomes, survival rates, and complication rates between coagulopathy and non-coagulopathy patients

Coagulopathy occurred in 26.9% of included patients. No significant differences in baseline characteristics were observed between coagulopathy patients and non-coagulopathy patients, except for gender (Table 1). As expected, APTT in coagulopathy patients was significantly prolonged when compared to non-coagulopathy patients (median, IQR: 42.9 s [34.2–84.5] vs. 28.3 s [25.5–34.3], P < 0.01). The initial FDP in coagulopathy patients was significantly higher than that in non-coagulopathy patients (114.8 μg/mL [92.5–168.3] vs. 30.9 μg/mL [14.2–50.3], P < 0.01).

Although no significant difference in favorable neurological outcome (34.4% vs. 50.6%, P = 0.15) and survival rates (56.3% vs. 72.4%, P = 0.12) was observed between coagulopathy and non-coagulopathy patients, these values were lower in coagulopathy patients. There was no difference between two groups in the complication rate (P = 0.85).

Comparison of baseline characteristics between the control (35.5–37 °C) and MTH (32–34 °C) groups in the patients with or without coagulopathy

No significant differences in patient characteristics were observed between the control and MTH groups among patients with or without coagulopathy (Table 2).

Comparison of neurological outcomes, survival rates and complication rates between the control and MTH groups in patients with or without coagulopathy

Among patients with coagulopathy, favorable neurological outcomes and survival rates were comparable between the control and MTH groups (33.3% vs. 35.0%, P = 1.00; 50.0% vs. 60.0%, P = 0.72) with no difference in complication rates (Table 3, left).

In patients without coagulopathy, the survival rate was significantly lower and the complication rate was significantly higher in the MTH group compared to the
control group (89.3% vs. 64.4%, \(P = 0.02\) and 0% vs. 20.3%, \(P < 0.01\), respectively; Table 3, right).

Comparison of APTT and FDP levels between the fever control and MTH groups in patients with or without coagulopathy
In patients with coagulopathy, there was no significant difference in APTT or FDP levels between the two groups at the time of admission; however, APTT was significantly prolonged in the MTH group compared to the control group on day 3 (\(P < 0.05\); Fig. 2).

In patients without coagulopathy, there was no significant difference in APTT or FDP levels between the two groups at the time of admission; however, FDP levels were significantly lower in the MTH group compared to the control group on day 1 (\(P < 0.05\); Fig. 2).

Comparison of platelet counts between the control and MTH groups in patients with coagulopathy
In patients with coagulopathy, there were no significant differences in platelet counts between the two groups at the time of admission to 1 day after rewarming (Additional file 1).

Comparison of ICP between the fever control and MTH groups in patients with or without coagulopathy
In patients with coagulopathy, the median ICP was 20 (10–46) mmHg, 21 (14–34) mmHg, and 19 (13–37) mmHg on day 1, day 3, and 1 day after rewarming, respectively. In patients without coagulopathy, the median ICP was 15 (10–22) mmHg, 15 (9–19) mmHg, and 21 (16–33) mmHg on day 1, day 3, and 1 day after rewarming, respectively. ICP was significantly higher in patients with coagulopathy compared to patients without coagulopathy on day 3 (\(P < 0.01\)).

In patients with coagulopathy, ICP did not differ between the control and MTH groups on day 1, day 3, or 1 day after rewarming (Fig. 3, left). In patients without coagulopathy, ICP was significantly lower in the MTH group compared to the control group on day 1 and at 1 day after rewarming (Fig. 3, right).

### Table 1 Patient characteristics

|                      | Coagulopathy n = 32 | Non-coagulopathy n = 87 | \(P\)-value |
|----------------------|----------------------|--------------------------|-------------|
| Age (years)          | 42 (22–56)           | 42 (21–55)               | 0.55        |
| Male                 | 27 (87.1)            | 52 (61.1)                | < 0.01      |
| Systolic blood pressure (mmHg) | 151 (124–170)       | 142 (114–179)            | 0.48        |
| Heart rate (beats/min) | 83 (71–110)          | 86 (70–106)              | 0.89        |
| Glasgow Coma Scale score | 6 (4–7)             | 6 (4–7)                  | 0.60        |
| Unreactive pupil or pupils | 14 (43.8)           | 37 (42.5)                | 1.00        |
| Platelet counts (×10⁴/mm³) | 23.7 (18.4–29.3)   | 22.7 (17.1–27.4)         | 0.52        |
| APTT (s)             | 42.9 (34.2–84.5)     | 28.3 (25.5–34.3)         | < 0.01      |
| FDP (μg/mL)          | 114.8 (92.5–168.3)   | 30.9 (14.2–50.3)         | < 0.01      |
| Fibrinogen (mg/dL)   | 206 (140–267)        | 202 (165–243)            | 0.87        |
| TCDB classification  |                      |                          | 0.77        |
| Diffuse injury grade I | 0 (0)                | 2 (2.3)                  |             |
| Diffuse injury grade II | 11 (34.4)           | 25 (28.7)                |             |
| Diffuse injury grade III | 7 (21.9)            | 12 (13.8)                |             |
| Diffuse injury grade IV | 1 (3.1)             | 3 (3.5)                  |             |
| Non-evacuated mass/Evacuated mass | 1/12                | 4/41                     | 0.68        |
| Surgical operation for TBI | 16 (51.6)          | 58 (66.7)                | 0.19        |
| Injury severity score | 25 (21–34)          | 25 (17–30)               | 0.09        |
| AIS score for head   | 4 (4–5)              | 4 (4–5)                  | 0.16        |
| Favorable outcome    | 11 (34.4)            | 44 (50.6)                | 0.15        |
| Survival rate        | 18 (56.3)            | 63 (72.4)                | 0.12        |
| Overall complication rate | 4 (12.5)           | 12 (13.8)                | 0.85        |

Values are presented as n (%) or median (interquartile ranges, IQRs)

MTH mild therapeutic hypothermia, AIS abbreviated injury score, TBI traumatic brain injury, CT computed tomography, FDP fibrin degradation products, TCDB Traumatic Coma Data Bank
In the present post hoc study, coagulopathy occurred in 32 (26.9%) of 119 patients with severe TBI. In patients with coagulopathy, favorable neurological outcomes were recorded in one third of patients and their survival rate was greater than 50%. Outcomes were similar between the MTH and control groups with no significant difference in complication rate, although prolongation of APTT lasted to day 3 in the MTH group. Consequently, we posit that both control and MTH have utility in the treatment of severe TBI in patients with coagulopathy on admission.

Tokutomi et al. examined the effects of hypothermia on several coagulation parameters (PT, APTT, platelet count, and AT-III) and performed a comparison with normothermia in TBI patients. They reported a trend toward a prolonged APTT in the hypothermia group on days 5 and 7 ($P = 0.07$ and $P = 0.06$, respectively).

**Table 2** Comparison of patient characteristics

| Variable                  | Coagulopathy | Non-Coagulopathy |
|---------------------------|--------------|------------------|
|                           | Control (35.5–37.0 °C) | MTH (32.0–34.0 °C) | P-value | Control (35.5–37.0 °C) | MTH (32.0–34.0 °C) | P-value |
|                           | n = 12       | n = 20           |         | n = 28       | n = 59           |         |
| Age (years)               | 31 (21–55)   | 48 (25–58)       | 0.34    | 41 (23–57)   | 42 (20–55)       | 0.93    |
| Male                      | 9 (81.8)     | 18 (90.0)        | 0.60    | 16 (59.3)    | 36 (62.1)        | 0.82    |
| Systolic blood pressure (mmHg) | 162 (128–177) | 138 (122–160)   | 0.15    | 142 (115–183)| 143 (110–175)   | 0.88    |
| Heart rate (beats/min)    | 77 (71–95)   | 94 (71–113)      | 0.34    | 86 (65–106)  | 85 (72–104)      | 0.94    |
| Glasgow Coma Scale score  | 6 (5–7)      | 6 (4–7)          | 0.72    | 6 (5–7)      | 6 (4–7)          | 0.65    |
| Unreactive pupil or pupils| 5 (41.7)     | 9 (45.0)         | 1.00    | 12 (42.9)    | 25 (42.4)        | 1.00    |
| Platelet counts ($\times 10^4$/mm$^3$) | 22.4 (18.7–29.3) | 24.5 (16.8–29.2) | 0.91    | 26.2 (16.7–29.8)| 22.1 (17.1–26.2) | 0.14    |
| APTT (s)                  | 42.8 (27.3–81.9) | 44.9 (35.4–87.1) | 0.73    | 27.4 (24.3–36.2)| 289 (25.7–33.2) | 0.58    |
| FDP ($\mu$g/mL)           | 168 (88.1–200.5) | 106 (92.4–142.2) | 0.17    | 37.5 (12.3–47.9) | 26 (16.5–55.6) | 0.91    |
| Fibrinogen (mg/dL)        | 228 (125–271) | 189 (164–246)    | 0.94    | 203 (167–264) | 199 (164–235)    | 0.49    |
| TCDB classification       |              |                  | 0.77    |              |                  | 0.30    |
| Diffuse injury grade I    | 0 (0)        | 0 (0)            | 1.00    | 1 (3.6)      | 1 (1.7)          |         |
| Diffuse injury grade II   | 5 (41.7)     | 6 (30.0)         | 8 (28.6) | 17 (28)     |                |         |
| Diffuse injury grade III  | 2 (16.7)     | 5 (25.0)         | 6 (21.4) | 6 (10.2)    |                |         |
| Diffuse injury grade IV   | 0 (0)        | 1 (5.0)          | 2 (7.1)  | 1 (1.7)     |                |         |
| Non-evacuated mass/Evacuated mass | 0/5 | 1/7 | 0.81 | 0/11 | 4/30 | 0.56 |
| Surgical operation for TBI| 7 (58.3)     | 9 (47.4)         | 0.72    | 18 (64.3)    | 40 (67.8)        | 0.81    |
| Injury severity score     | 25 (21–25)   | 34 (22–36)       | 0.08    | 22 (16–29)  | 25 (17–34)       | 0.36    |
| AIS score for head        | 5 (4–5)      | 4 (4–5)          | 0.50    | 4 (4–5)      | 4(4–5)           | 0.20    |
| AIS score ≥4 for other organs | 0 (0) | 4 (20.0) | 0.27 | 3 (10.7) | 4 (6.8) | 0.67 |

Values are presented as n (%) or median (interquartile ranges, IQRs) in the text.

**Table 3** Comparison of neurological outcomes and complication rates between coagulopathic and non-coagulopathic patients, and between the MTH (32–34 °C) and control (35.5–37 °C) groups with or without coagulopathy

| Variable                  | Coagulopathy | Non-Coagulopathy |
|---------------------------|--------------|------------------|
|                           | Control (35.5–37.0 °C) | MTH (32.0–34.0 °C) | P-value | Control (35.5–37.0 °C) | MTH (32.0–34.0 °C) | P-value |
|                           | n = 12       | n = 20           |         | n = 28       | n = 59           |         |
| Favorable outcome         | 4 (33.3)     | 7 (35.0)         | 1.00    | 16 (57.1)    | 28 (47.5)        | 0.49    |
| Survival rate             | 6 (50.0)     | 12 (60.0)        | 0.72    | 25 (89.3)    | 38 (64.4)        | 0.02    |
| Overall complication rate | 1 (8.3)      | 3 (15.0)         | 1.00    | 0 (0)        | 12 (20.3)        | < 0.01  |

MTH mild therapeutic hypothermia
Neurological outcomes were evaluated 6 months after brain injury
Complications occurring during targeted temperature management were recorded

**Discussion**

In the present post hoc study, coagulopathy occurred in 32 (26.9%) of 119 patients with severe TBI. In patients with coagulopathy, favorable neurological outcomes were recorded in one third of patients and their survival rate was greater than 50%. Outcomes were similar between the MTH and control groups with no significant difference in complication rate, although prolongation of APTT lasted to day 3 in the MTH group. Consequently, we posit that both control and MTH have utility in the treatment of severe TBI in patients with coagulopathy on admission.

Tokutomi et al. examined the effects of hypothermia on several coagulation parameters (PT, APTT, platelet count, and AT-III) and performed a comparison with normothermia in TBI patients. They reported a trend toward a prolonged APTT in the hypothermia group on days 5 and 7 ($P = 0.07$ and $P = 0.06$, respectively).
present study, the APTT in the MTH group at the time of admission (28.8 ± 2.7 s) did not differ from that of the control group (29.3 ± 5.7 s). Although the inclusion criterion used in the present study was a severely prolonged APTT of >60 s [23], the APTT in the MTH group among patients with coagulopathy was comparable to Tokutomi’s data and found to be significantly prolonged on day 3 compared to the control group [23].

Genet et al. examined the pathophysiology of trauma-induced coagulopathy between isolated TBI and non-TBI patients, and concluded that hemostatic, vascular, and endothelial responses were comparable [24]. Theoretically, APTT indicates the activity of the intrinsic and common pathways of coagulation, and prolongation of the APTT indicates the presence of a coagulation disorder [25]. Among patients with coagulopathy included in the present study, APTT was prolonged in the MTH group compared to the fever control group and reached statistical significance on day 3 (Fig. 2a). These findings are attributable to the speed of decreased biochemical reactions due to the decreased core body temperature in the MTH group. A similar trend was observed in patients without coagulopathy; however, this difference did not reach statistical significance (Fig. 2b). On the other hand, FDP levels in patients with coagulopathy were markedly elevated at the time of admission, as expected (Fig. 2c). The elevation

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**Fig. 2** Alterations of APTT and FDP between the fever control and MTH groups in patients with or without coagulopathy. 

- **a** Comparisons of APTT between the control and the MTH groups in the patients with coagulopathy. 
- **b** Comparisons of APTT between the control and the MTH groups in the patients without coagulopathy. 
- **c** Comparisons of FDP between the control and the MTH groups in the patients with coagulopathy. 
- **d** Comparisons of FDP between the control and the MTH groups in the patients without coagulopathy. The control group (35.5 °C–37 °C) is indicated in white and the MTH group (32 °C–34 °C) is indicated in gray. The boxes are the 25th to 75th percentiles and the whiskers are 5th to 95th percentiles. *p < 0.05; statistically significant. APTT: activated partial thromboplastin time, FDP: fibrin/fibrinogen degradation products.

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**Fig. 3** Alterations of ICP between the control and MTH groups in patients with or without coagulopathy. 

- **a** Comparisons of ICP between the control and the MTH groups in the patients with coagulopathy. 
- **b** Comparisons of ICP between the control and the MTH groups in the patients without coagulopathy. The control (35.5–37 °C) group is indicated in white and the MTH (32–34 °C) group is indicated in gray. The boxes are the 25th to 75th percentiles and the whiskers are 5th to 95th percentiles. *p < 0.05; statistically significant.
indicated acceleration of systemic fibrin deposition and secondary fibrinolysis/fibrinogenolysis by plasmin [24]. The values gradually decreased until day 3, both in the control and MTH groups, without statistical significance.

Polderman reported that very mild hypothermia (35 °C) does not affect coagulation and can be safely used even in patients at high risk of bleeding [9]. In addition, regarding mild hypothermia (33–35 °C), Wolberg et al. examined healthy volunteers and found that enzyme activities and platelet activation were not reduced at 33 °C [26]. Therefore, Gando et al. posited that isolated mild hypothermia at 33 to 35 °C does not have severe effects on hemostasis in typical clinical trauma settings [27]. Based on our study, MTH did not seem to negatively affect the outcomes in patients with coagulopathy following severe TBI on admission; therefore, the present study indicates that MTH may be applicable even in patients with severe TBI and coagulopathy. Typically, TBI-associated coagulopathy is not related to visual blood loss [28]; therefore, clinicians should attend to intracranial hemorrhage and organs with ongoing bleeding. In fact, patients with an AIS score ≥ 3 for other organs treated with mild therapeutic hypothermia had a mortality rate greater than 80% in the present study (data not shown).

In the patients with coagulopathy, ICP tended to be high with wide ranges both in the control and the MTH groups, compared to those in the patients without coagulopathy during the periods of the targeted temperature interventions, day 1 and day 3 (Fig. 3a, b). These differences might contribute their low rates of favorable neurological outcome and survival both in the fever control and MTH groups with coagulopathy (Table 3).

There are several limitations to the present study. First, the original study was terminated before the full sample size had been recruited. Additionally, the sample size was further reduced from 150 to 119 patients as APTT and/or FDP values could not be obtained in 16 out of 135 patients. These factors may have biased the outcomes of the present study. Second, d-dimer levels are the most specific test for coagulopathy in TBI [29]; however, the number of patients with d-dimer levels measured at the time of arrival was small in the present study. Further, PT could not be examined due to unavailability of the dataset. Third, an extremely small number of patients were included in this study (coagulopathy occurred in only 32 patients), and the results require confirmation in a larger cohort. Beta-error may have also existed. Fourth, although there was no significant difference in age (p = 0.34), patients in fever control group were younger than those in MTH group (median age 31 years vs. 48 years). Therefore, additional studies adjusted for background factors will be required. Finally, as the present study was a post hoc sub-analysis, selection bias may have been present.

Conclusion
Our study suggests that in comparison to control, MTH does not worsen the outcome of patients with coagulopathy following severe TBI.

Additional file

Additional file 1: Comparison of platelet counts between the control and MTH groups in patients with coagulopathy. MTH, mild therapeutic hypothermia. Values are presented as median (interquartile ranges, IQRs).

Abbreviations
AIS: Abbreviated Injury Severity Score; APTT: activated partial thromboplastin time; B-HYPO: Brain Hypothermia study group; FDP: fibrin/fibrinogen degradation products; GCS: Glasgow Coma Scale; ICP: intracranial pressure; ISS: Injury Severity Score; MTH: mild therapeutic hypothermia; RCT: randomized controlled trial; TBI: traumatic brain injury; TTM: targeted temperature management

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Not applicable.

Authors’ contributions
TH planned and conducted the statistical analyses on the database, appraised the background literature, prepared the first draft of the manuscript, and coordinated subsequent revisions. KK, SY, and KD contributed to collect the data. YK, YO and TM contributed to planning the analyses and to drafting and reviewing the manuscript. All authors read and approved the final manuscript and agree to be accountable for all aspects of the work.

Ethics approval and consent to participate
The protocol was approved by the Institutional Review Board of each participating hospital.

Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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References

1. Epstein DS, Mitra B, O'Reilly G, Rosenfeld JV, Cameron PA. Acute traumatic coagulopathy in the setting of isolated traumatic brain injury: a systematic review and meta-analysis. Injury. 2014;45(5):819–24.

2. Harhangi BS, Kompanjie EI, Leebeek FW, Maas AI. Coagulation disorders after traumatic brain injury. Acta Neurochir. 2008;150(2):165–75. discussion 175

3. Lustenberger T, Talving P, Kobayashi L, Barmparas G, Inaba K, Lam L, Branco BC, Demetriades D. Early coagulopathy after isolated severe traumatic brain injury: relationship with hypoperfusion challenged. J Trauma. 2010;69(6):1410–4.

4. Talving P, Benfield R, Hadjiaacharia P, Inaba K, Chan LS, Demetriades D. Coagulopathy in severe traumatic brain injury: a prospective study. J Trauma. 2009;66(5):61–6. discussion 61–52.

5. Wafaisade A, Lefering R, Tjardes T, Wutzler S, Simanski C, Paffrath T, Fischer P, Boullon B, Maegle M, Trauma Registry of DGU. Acute coagulopathy in isolated blunt traumatic brain injury. Neurocrit Care. 2010;12(2):211–9.

6. Chhabra G, Ranganathan K, Subramanian A, Agrawal D, Sharma S, Mukhopadhayakh AK. Hypofibrinogenemia in isolated traumatic brain injury in Indian patients. Neurol India. 2010;58(5):756–7.

7. Kearney TJ, Benet L, Grode M, Lee S, Hiat JR, Shabot MW. Coagulopathy and catecholamines in severe head trauma. J Trauma. 1992;32(5):608–11. discussion 610–2.

8. Endo A, Shihaishi A, Otomo Y, Kushimoto S, Saitsu D, Hayakawa M, Ogura H, Murata K, Hagiwara A, Sasajki J, et al. Development of novel criteria of the “lethal triad” as an indicator of decision making in current trauma care: a retrospective multicenter observational study in Japan. Crit Care Med. 2016;44(9):e977–803.

9. Polderman KH. Mechanisms of action, physiological effects, and complications of hypothermia. Crit Care Med. 2009;37(7 Suppl):S186–202.

10. Polderman KH, Herold I. Therapeutic hypothermia and controlled normothermia in the intensive care unit: practical considerations, side effects, and cooling methods. Crit Care Med. 2009;37(3):110–20.

11. Clifton GL, Miller ER, Choi SC, Levin HS, McCauley S, Smith KR Jr, Muizelaar JP, Wagner FC Jr, Marion DW, Luerksen TG, et al. Lack of effect of induction of hypothermia after acute brain injury. N Engl J Med. 2001;344(8):556–63.

12. Clifton GL, Valadka A, Zygour D, Coffey CS, Drever P, Fourwinds S, Janis LS, Wilde E, Taylor P, Harshman K, et al. Very early hypothermia induction in patients with severe brain injury (the National Acute Brain Injury Study: hypothermia II): a randomised controlled trial. Lancet Neurol. 2011;10(2):131–9.

13. Hutchison JS, Ward RE, Lacroix J, Hebert PC, Barnes MA, Bohn DJ, Dirks PB, Doucette S, Ferguson D, Gottesman R, et al. Hypothermia therapy after traumatic brain injury in children. N Engl J Med. 2006;355(23):2447–56.

14. Cariou A, Payen JF, Asehnoune K, Audibert G, Botte A, Brissaud O, Debaty G, Deltour S, Deye N, Engrand N et al. Therapeutic hypothermia versus fever control with tight hemodynamic monitoring and slow rewarming in patients with severe traumatic brain injury: a randomized controlled trial. J Neurotrauma. 2015;32(7):422–9.

15. Andrews PJ, Sinclair HL, Rodriguez A, Harris BA, Battison CG, Rhodes JK, Murray GD. Hypothermia for intracranial hypertension after traumatic brain injury. N Engl J Med. 2013;373(25):2403–12.

16. Chang EF, Meeker M, Holland MC. Acute traumatic intraparenchymal hemorrhage: risk factors for progression in the early post-injury period. Neurosurgery. 2006;58(4):647–9. discussion 647–56.

17. Brohi K, Cohen MJ, Davenport RA. Acute coagulopathy of trauma: mechanism, identification and effect. Curr Opin Crit Care. 2007;13(6):680–5.

18. Brohi K, Cohen MJ, Ganter MT, Matthay MA, Mackersie RC, Pittet JF. Acute traumatic coagulopathy: initiated by hyperperfusion: modulated through the protein C pathway? Ann Surg. 2007;245(5):812–8.

19. Brohi K, Singh J, Heron M, Coats T. Acute traumatic coagulopathy. J Trauma. 2003;54(6):1127–30.

20. Hifumi T, Krooda Y, Kawakita K, Yamashita S, Oda Y, Doi K, Maekawa T. Fever control management is preferable to mild therapeutic hypothermia in traumatic brain injury patients with abbreviated injury scale 3-4: a multi-center, randomized controlled trial. J Neurotrauma. 2016;33(11):1047–53.

21. Sawamura A, Gando S, Hayakawa M, Hoshino H, Kubota N, Sugano M. Effects of antithrombin III in patients with disseminated intravascular coagulation diagnosed by newly developed diagnostic criteria for critical illness. Clin Appl Thromb Hemost. 2009;15(5):561–6.

22. Tokutomi T, Miyagi T, Morimoto K, Karukaya T, Shigemori M. Effect of hypothermia on serum electrolyte, inflammation, coagulation, and nutritional parameters in patients with severe traumatic brain injury. Neurocrit Care. 2004;1(2):171–82.

23. Sawamura A, Hayakawa M, Gando S, Kubota N, Sugano M, Wada T, Katabami K. Disseminated intravascular coagulation with a fibrinolytic phenotype at an early phase of trauma predicts mortality. Thromb Res. 2009;124(5):608–13.

24. Poller L. Standardization of the APTT test. Current status. Scand J Haematol Suppl. 1980;37:49–63.

25. Wolberg AS, Meng ZH, Monroe DM 3rd, Hoffman M. A systematic evaluation of the effect of temperature on coagulation enzyme activity and platelet function. J Trauma. 2004;56(6):1221–8.

26. Gando S, Sawamura A, Hayakawa M. Trauma, shock, and disseminated intravascular coagulation: lessons from the classical literature. Ann Surg. 2011;254(1):10–9.

27. Sawamura A, Hayakawa M, Gando S, Kubota N, Sugano M, Wada T, Katabami K. Application of the Japanese Association for Acute Medicine disseminated intravascular coagulation diagnostic criteria for patients at an early phase of trauma. Thromb Res. 2009;124(6):706–10.

28. Zhang D, Gong S, Jin H, Wang J, Sheng P, Zou W, Dong Y, Hou L. Coagulation parameters and risk of progressive hemorrhagic injury after traumatic brain injury: a systematic review and meta-analysis. Biomed Res Int. 2015;2015:261825.