Impact of non-neurological complications in severe traumatic brain injury outcome

Luisa Corral1,2*, Casimiro F Javierre2, Josep L Ventura1, Pilar Marcos3, José I Herrero1 and Rafael Mañez1,4

Abstract

Introduction: Non-neurological complications in patients with severe traumatic brain injury (TBI) are frequent, worsening the prognosis, but the pathophysiology of systemic complications after TBI is unclear. The purpose of this study was to analyze non-neurological complications in patients with severe TBI admitted to the ICU, the impact of these complications on mortality, and their possible correlation with TBI severity.

Methods: An observational retrospective cohort study was conducted in one multidisciplinary ICU of a university hospital (35 beds); 224 consecutive adult patients with severe TBI (initial Glasgow Coma Scale (GCS) < 9) admitted to the ICU were included. Neurological and non-neurological variables were recorded.

Results: Sepsis occurred in 75% of patients, respiratory infections in 68%, hypotension in 44%, severe respiratory failure (arterial oxygen pressure/oxygen inspired fraction ratio (PaO2/FiO2) < 200) in 41% and acute kidney injury (AKI) in 8%. The multivariate analysis showed that Glasgow Outcome Score (GOS) at one year was independently associated with age, initial GCS 3 to 5, worst Traumatic Coma Data Bank (TCDB) first computed tomography (CT) scan and the presence of intracranial hypertension but not AKI. Hospital mortality was independently associated with initial GCS 3 to 5, worst TCDB first CT scan, the presence of intracranial hypertension and AKI. The presence of AKI regardless of GCS multiplied risk of death 6.17 times (95% confidence interval (CI): 1.37 to 27.78) (P < 0.02), while ICU hypotension increased the risk of death in patients with initial scores of 3 to 5 on the GCS 4.28 times (95% CI: 1.22 to 15.07) (P < 0.05).

Conclusions: Low initial GCS, worst first CT scan, intracranial hypertension and AKI determined hospital mortality in severe TBI patients. Besides the direct effect of low GCS on mortality, this neurological condition also is associated with ICU hypotension which increases hospital mortality among patients with severe TBI. These findings add to previous studies that showed that non-neurological complications increase the length of stay and morbidity in the ICU but do not increase mortality, with the exception of AKI and hypotension in low GCS (3 to 5).

Introduction

Severe Traumatic Brain Injury (TBI) mortality and morbidity is frequently caused by the neurological consequences of the brain injury [1,2]. Nevertheless, non-neurological complications are also frequent, and may be cardiovascular, respiratory, infectious and others [3-5]. An initial study by Zygun et al. showed that non-neurological organ dysfunction measured by Sequential Organ Failure Assessment (SOFA) was not associated with increased mortality in neurocritical patients with TBI or subarachnoid hemorrhage, despite the frequent finding of cardiopulmonary dysfunction [6]. However, a later study by the same group associated non-neurological organ dysfunction with worse outcome, reporting that cardiovascular and respiratory complications were the most common dysfunctions [5].

The pathophysiology of systemic complications after TBI is unclear. Complications may arise from the direct effect of the injury or as a result of the side effects of therapy [7,8]. Neurogenic causes such as the massive catecholamine and neuro-inflammatory response associated with brain injury may contribute to systemic complications. In addition, the intensive care management of TBI patients is mainly directed at neurological problems and may contribute to non-neurological complications [8].
A better knowledge of the incidence, causes and consequences of non-neurological complications in patients with severe TBI would help in their prevention, treatment and prognosis. The purpose of this study was to analyze the non-neurological complications presented by patients with severe TBI admitted to the ICU, the impact of these complications on hospital mortality and outcome at one year, and their possible correlation with TBI severity.

Materials and methods
The study included 224 adult patients with severe TBI (initial Glasgow Coma Scale (GCS) < 9), consecutively admitted to the ICU, between 1998 and 2004. It is an observational retrospective cohort study. The initial GCS score was obtained upon admission to the emergency ICU. If the patient was already intubated and on mechanical ventilation the previous GCS score was used and GCS scale was divided into two levels of severity: low (GCS 3 to 5) and high (GCS 6 to 8) [9,10]. The study was approved by the institution’s Research Ethics Committee, which did not consider informed consent to be necessary because of the study design. Management of patients followed international guidelines [11], with the goal of maintaining a cerebral perfusion pressure (CPP) of 60 to 70 mmHg and an intracranial pressure (ICP) of < 20 mmHg. Intracranial hypertension was defined as ICP > 20 mmHg for 15 minutes without a systemic cause. Elevations in ICP were managed sequentially with sedation, paralysis, mannitol and mild hyperventilation. Barbiturate therapy was considered for refractory intracranial hypertension. The therapy index [12] was used to quantify the treatment for intracranial hypertension that was enough to quantify the treatments used in that year following international guidelines. Computerized tomography (CT) of the brain at admission was evaluated following the Traumatic Coma Data Bank Computed Tomography (TCDB CT) scan classification. Mortality at hospital discharge and Glasgow Outcome Scale (GOS) at one year was obtained and was dichotomized as worse outcome (dead, vegetative state and severe disability) and as good outcome (moderate disability and good recovery) [9].

The following non-neurological complications were recorded:

Cardiovascular: ICU hypotension (systolic blood pressure (SBP) < 90 mmHg for 30 minutes during the ICU stay), hypertension (SBP > 160 mmHg for more than 30 minutes and, in hypertensive subjects, an increase > 40 mmHg that required treatment change [3]), cardiac arrhythmias (bradycardia < 60 beats/minute or tachycardia > 120 beats/minute) and the need for vasoactive drugs (dopamine, dobutamine or norepinephrine).

Respiratory: infectious respiratory complications (aspiration, tracheobronchitis or pneumonia (infiltrate on chest radiograph and analysis of gram or sputum culture-positive [3]), acute respiratory distress syndrome (ARDS) [13], PaO$_2$/FiO$_2$ ratio 200 to 300 and severe respiratory failure (SRF) as PaO$_2$/FiO$_2$ ratio < 200.

Septic: severe sepsis [14] and septic shock [14].

Renal: acute kidney injury (AKI) (serum creatinine > 150 μmol/L) with or without dialysis [3].

Abdominal/digestive complications: ileus, cholecystitis, bilirubin levels > 18 μmol/L and aspartate aminotransferase (AST) > 1 μkat/L [3].

Endocrinometabolic: hyponatremia < 130 mmol/L, hypernatremia > 150 mmol/L, syndrome of inappropriate antidiuretic hormone secretion (SIADH) (Na < 130 mmol/L, urine osmolality exceeding serum osmolality [3]), cerebral salt wasting syndrome (excessive diuresis with urine sodium below > 20 mmol/L [15]) and diabetes insipidus (diuresis > 200 mL/hour for 24 hours without response to fluid restriction or needing treatment with desmopressin, urine specific gravity < 1,005 and a urine osmolality less than half that of plasma).

Bleeding: hemorrhage needing blood transfusion products (> 4 packed cells) and/or hemorrhagic shock during ICU admission.

Statistical analysis
Continuous variables (age, ICU stay and in hospital stay) were expressed as median and interquartile range. Categorical data were expressed as frequency and percentage. We used the chi square test to compare categorical data and proportions, and the T-test, Mann-Whitney test or Kruskal-Wallis, as appropriate, to compare continuous variables. Multivariate analysis was performed with multivariate regression and variables were selected using either the stepwise or the backward regression procedures. Confounding and interaction variables were studied, and the effect size on outcome was calculated for clinical complications during the ICU stay, as well as the effect size on ICU hypotension. An alpha level of 0.05 was used to determine statistical significance. All data in the present study were analyzed using SPSS version 16.0 (SPSS Inc, Chicago, USA).

Results
The characteristics of the sample are shown in Table 1 [9,16]. ICP was monitored in 73% of the patients (n = 164), 51% (n = 83) of whom developed intracranial hypertension and 56% low cerebral perfusion pressure at some point in the evolution of their condition. Hypoxia at admission was detected in 64 patients, it was not detected in 152 and 8 had missing data. Hypotension at admission was detected in 38 patients. The median
therapy index score was 3, with 50% of the patients ranging between 2 and 7. The median ICU stay was 15 days (interquartile range 20).

Non-neurological complications throughout the ICU stay are shown in Table 2: sepsis occurred in 75% of the patients, respiratory infections in 68%, ICU hypotension in 44%, severe respiratory failure (PaO2/FiO2 < 200) in 41% and AKI in 8%. Vasoactive drugs were used in 96% of the patients with ICU hypotension. There was only one patient with previous chronic renal failure, which was not included as AKI. The univariate analysis of the non-neurological complications showed that hypotension, severe respiratory failure, septic shock, AKI, bleeding complications and non-neurological surgery were prognostic factors related to hospital mortality (Table 3). A multivariate regression analysis was performed for hospital mortality and GOS at one year post-TBI, taking into account variables with clinical relevance (age), variables related to TBI severity [(initial GCS (divided into GCS 3 to 5 and GCS 6 to 8), first CT scan (TCDB) and the presence of intracranial hypertension) and non-neurological complications that were statistically significant in the univariate analysis. The multivariate analysis showed that the worst outcome at one year was independently associated with age, initial GCS 3 to 5, worst TCDB first CT scan and the presence of intracranial hypertension but not AKI (Table 4). The multivariate analysis showed that hospital mortality was independently associated with an initial GSC 3 to 5, worst TCDB first CT scan, the presence of intracranial hypertension and AKI (Table 5). Neither multiple trauma nor chest trauma was independently associated with higher mortality or worse outcome. Seventeen patients (8%) developed AKI and 13 of them (76%) died. In 10 of the 13 patients (77%) death occurred after day 7 of ICU admission and in 4 of 13 renal replacement techniques had been used.

The specific effect of ICU hypotension, severe respiratory failure or AKI on outcome was also assessed in a multivariate regression model that included age, initial

### Table 1 Clinical characteristics of 224 patients with severe traumatic brain injury.

| Variables                      | No. (%) |
|--------------------------------|---------|
| **Age (median, interquartile range)** | 35.6 (23-55) |
| **Gender**                      |         |
| Male                            | 189 (84) |
| Female                          | 35 (16)  |
| **Mechanism of injury**         |         |
| Traffic accident                | 148 (66) |
| Falls                           | 58 (26)  |
| Assault                         | 6 (3)    |
| Others                          | 12 (5)   |
| **Multiple trauma**             |         |
|                                | 127 (57) |
| **GCS**                         |         |
| 3                               | 30 (13)  |
| 4                               | 35 (16)  |
| 5                               | 30 (13)  |
| 6                               | 40 (18)  |
| 7                               | 35 (16)  |
| 8                               | 54 (24)  |
| Low GCS (GCS 3, 4, 5)           | 95 (42)  |
| High GCS (GCS 6, 7, 8)          | 129 (58) |
| **TCDB CT scan classification** |         |
| I                               | 14 (6)   |
| II                              | 120 (54) |
| III                             | 26 (12)  |
| IV                              | 21 (9)   |
| EM                              | 38 (17)  |
| NEM                             | 5 (2)    |
| I, II, EM                       | 172 (77) |
| III, IV, NEM                    | 52 (23)  |
| **Detected hypotension at admission** | 38 (17) |
| **Detected hypoxia at admission** | 64 (29) |
| **Detected intracranial hypertension** | 83 (37) |
| **Pupil abnormalities**         | 76 (34)  |
| **Dead**                        |         |
| At ICU discharge                | 67 (31)  |
| At hospital discharge           | 74 (33)  |
| At 6 and 12 months              | 79 (35)  |
| **Days in ICU (median, interquartile range)** | 15 (7-26) |
| **Days in hospital (median, interquartile range)** | 26 (11-49) |

EM: evacuable mass, GCS: Glasgow Coma Scale, ICU: Intensive Care Unit, NEM: non-evacuable mass, TCDB CT: Traumatic Coma Data Bank Computed tomography.

TCDB CT was grouped in two groups: one best TCDB CT scan classification type I, II and evacuable mass (EM) and the other worst TCDB CT scan type III, IV and non-evacuable mass (NEM).

### Table 2 Non-neurological complications.

| Variables                      | n (%) |
|--------------------------------|-------|
| Respiratory infections         | 152 (68) |
| Atelectasis                    | 47 (21) |
| ARDS                           | 20 (9)  |
| PaO2/FiO2 < 200                | 92 (41) |
| PaO2/FiO2 200-300              | 94 (42) |
| Cardiovascular                 |       |
| Hypotension                    | 99 (44) |
| Hypertension                   | 28 (12) |
| Arrhythmias                    | 25 (11) |
| Dopamine, dobutamine or norepinephrine | 156 (70) |
| Infection                      |       |
| Sepsis                         | 169 (75) |
| Septic shock                   | 13 (6)  |
| AKI                            | 17 (8)  |
| Abdominal complications        | 40 (18) |
| Electrolytical complications   | 48 (21) |
| Bleeding complications         | 60 (27) |
| Non-neurosurgical surgery      | 62 (28) |

AKI: acute kidney injury, ARDS: acute respiratory distress syndrome, PaO2/FiO2: arterial oxygen pressure/oxygen inspired fraction ratio.

Respiratory infections: aspiration, tracheobronchitis and pneumonia were grouped. Non-neurosurgical surgery: traumatic, digestive, maxillar or plastic surgery.
GCS, initial CT scan (TCDB), the presence of intracranial hypertension, the two other non-neurological complications (sepsis and respiratory infections) and their interactions. The presence of AKI regardless of GCS multiplied the risk of death 6.17 times with 95% CI of 1.37 to 27.78 ($P < 0.02$) and ICU hypotension in patients with low initial GCS (3 to 5) significantly increased the risk of death (4.28 times with 95% CI of 1.37 to 27.78, $P < 0.05$) (Table 6).

To estimate the effect of initial GCS on the development of non-neurological complications a multivariate regression analyzed the interactions between these conditions, finding a single interaction, low initial GCS (3 to 5), multiplied the risk of developing ICU hypotension 3.37 times ($P < 0.005$) (Table 6), suggesting that the latter condition was related to the neurological situation.

Finally, we also observed the impact of non-neurological complications in the ICU stay of surviving patients with severe TBI. The ICU stay (median (interquartile range)) in surviving patients was longer when hypotension (16 (17) versus 30 (27) days, $P < 0.01$), severe respiratory failure (16 (20) versus 25 (28) days, $P < 0.01$) and AKI (18 (20) versus 39 (39) days, $P < 0.05$) were present.

**Discussion**

The present study found that non-neurological complications were frequent during the ICU stay in a cohort of patients with severe TBI. We found some differences vis-à-vis other studies with regard to non-neurological conditions leading to worse prognosis. Hypotension, pneumonia, infectious complications and coagulation dysfunction have been associated with an unfavorable outcomes [1,2,4,17,18] but in our study only AKI and hypotension in low GCS contributed independently to increasing hospital mortality together with the TBI severity variables (low initial GCS, first worst CT and intracranial hypertension).

AKI has been considered an uncommon non-neurological complication of severe TBI and is not associated

### Table 3 Non-neurological complications associated to hospital mortality.

| Variables                | Deaths 74 (33%) | Survive 150 (67%) | $P$  |
|--------------------------|-----------------|-------------------|------|
|                          | n (%)           | n (%)             |      |
| Respiratory              |                 |                   |      |
| Respiratory infection    | 45 (30)         | 107 (70)          | NS   |
| Atelectasis              | 11(23)          | 36 (77)           | NS   |
| ARDS                     | 18 (90)         | 2 (10)            | < 0.01 |
| PaO$_2$/FiO$_2$ < 200    | 43 (47)         | 49 (53)           | < 0.05 |
| PaO$_2$/FiO$_2$ 200-300  | 21 (22)         | 73 (78)           | < 0.05 |
| Cardiovascular           |                 |                   |      |
| Hypotension              | 50 (51)         | 49 (49)           | < 0.05 |
| Hypertension             | 7 (25)          | 21 (75)           | NS   |
| Arrhythmias              | 12 (48)         | 13 (52)           | NS   |
| Dopamine, dobutamine or norepinephrine | 63 (40) | 93 (60) | < 0.05 |
| Infection                |                 |                   |      |
| Sepsis                   | 53 (31)         | 116 (69)          | NS   |
| Septic shock             | 11 (85)         | 2 (15)            | < 0.05 |
| AKI                      | 13 (76)         | 4 (24)            | < 0.05 |
| Abdominal complications   | 32 (30)         | 76 (70)           | NS   |
| Electrolytical complications | 18 (38)   | 29 (62)           | NS   |
| Bleeding complications    | 30 (50)         | 30 (50)           | < 0.05 |
| Non-neurosurgical surgery | 13 (21)        | 48 (79)           | < 0.05 |

AKI: acute kidney injury, ARDS: acute respiratory distress syndrome, PaO$_2$/FiO$_2$: arterial oxygen pressure/oxygen inspired fraction ratio. Respiratory infections: aspiration, tracheobronchitis and pneumonia were grouped.

### Table 4 Logistic regression backward stepwise for worse outcome at one year.

| Variable                  | OR   | IC 95%     | $P$ value |
|---------------------------|------|------------|-----------|
| Age                       | 1.03 | 1.01-1.06  | 0.007     |
| GCS 3-5                   | 3.26 | 1.36-7.89  | 0.008     |
| Worst TCDB first CT scan  | 3.05 | 1.15-8.12  | 0.025     |
| Intracranial hypertension | 3.83 | 1.72-8.53  | 0.001     |

GCS: Glasgow Coma Scale, IC 95%: 95% confidence interval for the OR, OR: odds ratio, TCDB CT: traumatic coma data bank computed tomography.

Worst TCDB CT scan referring to type III, IV and non-evacuable mass.

### Table 5 Logistic regression backward stepwise for mortality.

| Variable                  | OR   | IC 95%     | $P$ value |
|---------------------------|------|------------|-----------|
| GCS 3-5                   | 2.47 | 1.03-5.93  | 0.043     |
| Worst TCDB first CT scan  | 3.69 | 1.43-9.51  | 0.007     |
| Intracranial hypertension | 10.18| 3.85-26.89 | 0.000     |
| Renal failure             | 7.12 | 1.58-31.98 | 0.010     |

GCS: Glasgow Coma Scale, IC 95%: 95% confidence interval for the OR, OR: odds ratio, TCDB CT: traumatic coma data bank computed tomography.

Worst TCDB CT scan referring to type III, IV and non-evacuable mass.
Table 6 Adjusted odds ratio for predictors of mortality and adjusted odds ratio for predictors of ICU hypotension in our cohort.

| Variable                        | OR   | 95% CI     | P value |
|---------------------------------|------|------------|---------|
| Mortality                       |      |            |         |
| ICU hypotension with low GCS    | 4.28 | 1.22-15.07 | 0.024   |
| GCS 3-5                         |      |            |         |
| ICU hypotension with high GCS   | 0.13 | 0.01-0.233 | 0.167   |
| GCS 6-8                         |      |            |         |
| Severe respiratory failure      | 1.39 | 0.57-03.35 | 0.467   |
| Renal failure                   | 6.17 | 1.37-27.78 | 0.018   |
| ICU hypotension                 | 3.37 | 1.66-6.82  | 0.001   |

GCS: Glasgow Coma Scale, IC 95%: 95% confidence interval for the OR, ICU: intensive care unit, OR: odds ratio.

Hypotension, both at admission and during the ICU stay, has been associated with poor prognosis in patients with severe TBI [17,18], so that we analyzed accurately. Duration of hypotension has also been considered one of the worst prognostic factors in these patients [1,18]. In our study, hypotension at some point during the ICU stay was frequent (44%). However, in contrast to other studies [3,5], it was not found to be an independent risk factor for mortality, although it increased mortality in patients with low GCS. Patients with low GCS (3 to 5) show global brain, brainstem or reticular system damage [23], which may involve the dysfunction of systemic homeostasis. This leads to non-neurological neurogenic dependent complications, which are difficult to distinguish because of the multiple conditions that these patients present, including neurogenic myocardial dysfunction and neurogenic pulmonary edema. The etiological hypothesis is that catecholamine release leads to these cardiopulmonary abnormalities. Dysregulated inflammatory mechanisms also appear to play a role in the development of the multiple organ dysfunction syndrome [5]. In our study, neurogenic causes appear to cause the hypotension that increases mortality in patients with low GCS. Despite using vasoactive drugs, a more aggressive resuscitation of patients who are at risk of hypotension, to improve outcome, requires further investigation. The bad neurological prognosis of these patients may be a limitation to using more aggressive life-sustaining therapies during their ICU stay.

Other complications, such as respiratory problems, were also frequently observed in patients with severe TBI in this study. Pneumonia was common (42%), as in other studies [3,4,24], but was not associated with increased mortality. Ventilator-associated pneumonia does not appear to raise in-hospital mortality in patients with severe TBI, but it prolongs mechanical ventilation, ICU and hospital length of stay, and increases the need for tracheostomy [25-27]. Severe respiratory failure occurred in 41% of patients, similar to another study [5]. Moderate and severe respiratory failure are poor prognostic factors [28-30], but in other studies they have not been associated with increased hospital mortality [19]. In the study by Zygun et al. neither SOFA-defined nor MOD-defined respiratory failure was significantly associated with increased hospital mortality [19]. In the study by Bratton et al. acute lung injury (ALI), PaO2/FiO2 < 300 was associated with the global severity of head injury (lower GCS scores) but not with specific anatomic lesions diagnosed by cranial CT scans [28]. In our study, severe respiratory failure was not independently associated with mortality, but it increased ICU stay. The measures proposed to prevent respiratory complications are: prevention of lung collapse and consolidation, prevention of lung infection, and acceleration of weaning from mechanical ventilation in the ICU [31]. The use of prone position, positive end-expiratory pressure (PEEP) and recruitment maneuvers have proved effective in improving tissue oxygenation with minimal negative effects of intracranial pressure and cerebral blood flow. Accurate monitoring of the physiologic cerebral function is warranted to minimize the possible negative effects on the brain induced by the maneuvers themselves [31-33]. Recently analyzed ventilator care...
bundles may be useful to prevent ventilator-associated pneumonia [34].

Intracranial hypertension is a poor prognostic factor in patients with severe TBI [1,18,35-39]. The presence of multiple trauma was not associated with mortality in this study. SOFA and MOD scales have been used to analyze non-neurological complications [5,6,19]. In one study, intracranial hypertension had higher maximum SOFA and delta SOFA, and there was a tendency for lower GCS to present higher maximum SOFA and delta SOFA which was not subsequently confirmed with the assessment of MOD [6]. However, the same authors found that the SOFA scoring system had superior discriminative ability and a stronger association with outcome than MOD with regard to hospital mortality and unfavorable neurological outcome [19]. In another study, patients with acute brain injury with GCS < 9 had a greater incidence of sepsis, respiratory failure and need for vasopressor therapy than those with GCS > 8 [40]. However, the relationship between non-neurological organ dysfunction and TBI severity has not always been demonstrated [7]. In our study, mortality depended mainly on the TBI initial severity (low GCS 3 to 5 and worst CT scan); as in previous studies, cardiovascular, respiratory and infectious problems were the most frequent non-neurological complications [3-5,19,40], but only AKI and hypotension in the low GCS group were independently associated with higher mortality.

This study has some limitations. This study was done in a single center. The definitions of non-neurological complications are not standardized; although a recent study tried to standardize data collection in TBI they did not define non-neurological complications [41]. The complications were associated with mortality, but each insult was not collected individually, so the particular impact of each one in secondary brain damage could not be evaluated. The episodes of hypotension were recorded from the first day of admittance in the ICU until the day of discharge, therefore the episodes could have appeared at any time during the period between admittance and discharge of the patients. Nor were the severity and timing of non-neurological complications recorded. However, it has been shown previously that time to development of these complications does not influence the mortality of patients with severe TBI [5], but the knowledge of the timing of these complications would be helpful to determine if these complications occurred in a pre-terminal phase or not. Withdrawal of life-sustaining therapies was not recorded.

Conclusions

In summary, low initial GCS, first CT scan and intracranial hypertension are known neurological variables that determine mortality in severe TBI patients. AKI was the only non-neurological complication associated with ICU mortality. These findings add to previous studies that showed that non-neurological complications increase length of stay and morbidity in the ICU, but do not increase mortality, with the exception of AKI and hypertension in low GCS (3 to 5).

Key messages

- GCS 3 to 5, first CT scan and intracranial hypertension are neurological variables that determine mortality in severe TBI patients.
- AKI and the association of low GCS with hypotension were the non-neurological complications independently associated with ICU mortality.
- Non-neurological complications increase length of stay and morbidity in ICU, but do not increase mortality.

Abbreviations

AKI: acute kidney injury; ALI: acute lung injury; APACHE: Acute Physiology and Chronic Health Evaluation; ARDS: Acute Respiratory Distress Syndrome; AST: aspartate amino transferase; CI: confidence interval; CPP: cerebral perfusion pressure; CT: computed tomography; GCS: Glasgow Coma Scale; GOOS: Glasgow Outcome Scale; ICP: intracranial pressure; MOD: multiple organ dysfunction; PEEP: positive end-expiratory pressure; RIFLE: Risk: Injury: Failure: Loss: and End-stage Kidney; SBP: systolic blood pressure; SIADH: Syndrome of Inappropriate Antidiuretic Hormone Secretion; SOFA: Sequential Organ Failure Assessment; SRF: severe respiratory failure; TBI: traumatic brain injury; TCDB: Traumatic Coma Data Bank.

Author details

1Intensive Care Unit, Hospital Universitari de Bellvitge, Feixa Llarga s/n, L’Hospitalet de Llobregat-08907, Barcelona, Spain. 2Department de Ciències Fisiològiques II, Universitat de Barcelona, Feixa Llarga s/n, L’Hospitalet de Llobregat-08907, Barcelona, Spain. 3Intensive Care Unit, Hospital Universitari Germans Trias i Pujol, Carretera de Canyet s/n, Badalona-08916, Barcelona, Spain. 4Institut d’Investigació Biomèdica de Bellvitge (IDIBELL), Feixa Llarga s/n, L’Hospitalet de Llobregat-08907, Barcelona, Spain.

Authors’ contributions

LC and JV conceived of the study and participated in its design. LC, JV and LH collected data. PM and CJ performed the statistical analysis. RM participated in the study coordination and helped to draft the manuscript. All authors read and approved the final manuscript.

Competing interests

The authors declare that they have no competing interests.

Received: 1 October 2011 Revised: 15 November 2011 Accepted: 12 March 2012 Published: 12 March 2012

References

1. Andrews PJ, Sleeman DH, Statham PF, McQuatt A, Crouble V, Jones PA, Howells TP, McMillan CS: Predicting recovery in patients suffering from traumatic brain injury by using admission variables and physiological data: a comparison between decision tree analysis and logistic regression. J Neurosurg 2002, 97:326-336.
2. Perel P, Arango M, Clayton T, Edwards P, Komolafe E, Pocock S, Roberts I, Shakur H, Steyerberg E, Yuthavongsut S: Predicting outcome after traumatic brain injury: practical prognostic models based on large cohort of international patients. BMJ 2008, 336:425-429.
3. Piek J, Chesnut RM, Marshall LF, van Berkum-Clark M, Klauber MR, Blunt BA, Eisenberg HM, Jane JA, Marmarou A, Foulines MA: Extracranial complications of severe head injury. J Neurosurg 1992, 77:901-907.
