The initial trauma associated hemorrhagic shock in the multiply injured patient has pro-inflammatory effects. A retrospective single center cohort study

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Abstract

Background: The development of a Systemic Inflammatory Response Syndrome (SIRS) has been discussed to be related to the trauma load. Trauma, especially the multiple injured patient suffers blood loss in a various extent. The outcome of such patients is also linked with the need for transfusions and SIRS. Therefore, the association between the initial blood loss and the incidence of IL6 release following multiple injuries was examined.

Methods: Totally 545 patients ≥16 years and with an ISS >16 were included into this retrospective cohort study. The sample was divided according to the ATLS shock classes I-IV on admission in the trauma bay and the shock-index (< 0.5; 0.5-1.5 and > 1.5) at trauma site. The systemic levels of IL6 were measured 24 and 48 hours after admission. Data were analyzed using Kruskal-Wallis H test. The predictive quality was tested using receiver-operating curves (AUC), the independent predictive power was analyzed using binary logistic regression. The analysis was conducted using IBM SPSS® 23.0. Level of evidence III.

Results: IL6 significantly increased within the first 24h according to shock (252ng – 872ng, p < 0.001). There were highly predictive values between IL6 and SIRS (24h) (AUC: 0.345 – 0.951 for ATLS shock I-IV, p < 0.001; AUC: 0.443 – 0.963 for shock-index 0.5 - 1.5, p < 0.001). ATLS shock revealed as an independent predictor for IL6 increase after 48h (p < 0.001).

Conclusion: Initial shock-index at trauma site and the ATLS shock degree in the trauma bay are high predictors for IL6 release what may serve as a primer for the induction of SIRS. Very early control of blood loss seems to be the key point. However, these results might also point on the change of transfusion protocols in the multiple injured patient.

Introduction

The physiological reaction to severe multiple trauma has been shown to result in the systemic inflammatory response syndrome (SIRS) which appears to be related to trauma severity [1-3]. A sustained systemic inflammation is associated with increased mortality in multiple injured patients and results in the compensatory anti-inflammatory response syndrome (CARS) as an initially parallel to SIRS running compensatory part with an immunological burn out and higher susceptibility to infections [3]. There were several trials to determine the origin or the trigger of SIRS during the last decades, though the studies remain descriptive and the initial trigger was not detected [4]. Among the predictors of SIRS, direct indicators such as injury severity (ISS, NISS) and indirect indicators such as the rate of blood transfusions were discussed [5-8]. Certainly, trauma load, blood loss, and amount of transfused blood go hand in hand and are a result of the trauma. IL6 is a pro-inflammatory cytokine secreted by plenty of cells mostly hematopoietic cells including tumor cells maintaining their survival by the JAK/STAT pathway [9]. The IL6 concentration in patient’s peripheral blood might mirror the possibility to develop SIRS and multi organ dysfunction in the case of multiple injuries [10]. This retrospective cohort study was designed on the question, whether the initial blood loss at trauma site and then in the trauma bay influences the pro-inflammatory state in the multiple injured patient. The fact is that trauma leads to blood loss and the multiple injured patients bleed in a physiologically relevant way. The question was whether the blood loss classified on the trauma scene by the shock index and in the trauma bay by the ATLS shock degree has a pro-inflammatory effect and an effect on the expression of IL6 in multiple injured patient arising a therapeutic consequence. Here the statistical proof is provided that IL6 is raised by the blood loss in the multiple injured patient.

Methods

Patient sample

In this retrospective cohort study, 545 patients with multiple trauma, admitted to the trauma bay of the University Hospital Zurich (Switzerland), were included. Totally, 2995 datasets had to be excluded due to missing IL6 data. The data were included 01.01.2000 – 12.31.2014.

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In this period, only 545 multiple injured patients with IL6 data were available. The inclusion criteria were an injury severity score (ISS) >16 points, age ≥16 years, and immediate admission to the trauma bay of the University Hospital Zurich after incurring the multiple trauma. The patient sample was divided into four groups according to the ATLS shock classes I-IV and the shock index < 0.5, 0.5-1.5 and > 1.5 [11].

Data collection

All data were collected retrospectively according to Good Clinical Practice Guidelines and were pooled in the Zurich Trauma Registry data bank. The patients' data were retrieved from patient records over an time period of 21 days, with the approval of the local institutional review board (IRB), according to the University of Zurich IRB guidelines and the World Medical Association Declaration of Helsinki. The study was conducted according to our institutional guidelines for good clinical practice (Ethics Committee of the University Hospital Zurich: "Retrospektive Analysen in der Chirurgischen Intensivmedizin" Nr. St.V. 01-2008). Level of Evidence III.

Scoring systems

The scoring systems were used to depict a detailed view on the multiple injured patient sample. The overall physiological impairment was evaluated by the acute physiology and chronic health evaluation (APACHE II) score of the patient at admission [12]. The ISS and the new injury severity scale (NISS) were used to define the severity of trauma [13,14]. The abbreviated injury scale (AIS; update 2008 version) was used to describe injuries in specific anatomical regions. The Trauma Score - Injury Severity Score (TRISS) was used to analyze the probability of death in the patients sample at admission [15]. The Simplified Acute Physiology Score (SAPS II) was calculated according to Le Gall at admission [16]. SIRS was defined as previously published and sepsis was defined as two or more SIRS criteria with an infectious focus [17].

Groups

To determine and to distinguish the Shock states two different staging methods were chosen and calculated retrospectively from patient's records for descriptive purposes.

1. The ATLS classification (Class I, II, III and IV) for severe hemorrhage was evaluated from the earliest available data at admission in the trauma bay.

2. The shock index according to Allgöwer was calculated from the earliest available data at trauma scene and subdivided into three groups [18]: <0.5, 0.5-1.5 and >1.5.

Laboratory parameters

Blood lactate levels, pH, and hematomas were measured at intervals using a blood gas analyzer (ABL800 Flex, Radiometer, Thalwil, Switzerland). The prothrombin time was measured using a standardized method [19].

Statistical analysis

Missing datasets were excluded. Data are presented as mean ± standard deviation for continuous variables and as percentages for categorical variables. The Two-tailed Kolmogorov-Smirnov test was used for normality testing if data is p > 0.05 than it was considered to be normally distributed. The data of the groups were compared using a χ² test for categorical data. The one-way analysis of variance (ANOVA) was taken for normally distributed continuous data. For not normally distributed continuous data, the Kruskal-Wallis H test was used. Results were considered significant if p < 0.05. If the p-value was 0.001 < p < 0.05 a post hoc analysis by Bonferroni was performed. Normality testing was done with the Kolmogorov-Smirnoff test, data were considered as normally distributed if p > 0.05. The predictive quality of the different injuries was reported as the area under the receiver operator characteristic curve (AUC). The independent predictive power was analyzed using binary logistic regression. The goodness of fit for the binary logistic regression was analyzed by the Hosmer-Lemeshow test and considered as good if p > 0.05. Data were analyzed using IBM SPSS® Statistics for Windows software (version 23.0; IBM Corp., Armonk, NY, USA).

Source of bias

The investigated effect might have many possible cofounders such as ISS, AIS, contamination, preexisting diseases and medication. However, these first data with such a high predictive quality and in this temporary context clearly point into this direction.

Results

Patient sample and the conditions at admission

There were significantly more men than women in each ATLS shock class group. The mean age was around forty years, without significant differences (Table 1). The ISS and NISS were significantly increasing along the ATLS shock classifications (Table 1). The physiological parameters, summarized in APACHE II, were rising along the shock severity as well as the according changes in Base Excess, Lactate, Hematocrit, Hemoglobin and Prothrombin Time (Table 1).

The IL 6 secretion pattern in Shock

The secretion pattern of IL 6 was similar in both distributions, the ATLS shock class and shock index (Figure 1A and B). In both distributions there was a significant increase found according to the initial blood loss. However, the performed post-hoc analysis revealed only significances in ATLS shock class IV vs. I, II and III after 24 hours. No significances were found after 48 hours. Post hoc analysis revealed a significant difference in ATLS shock class I vs. II after 24 hours. A similar statistical situation was found regarding the shock index distribution (Figure 1B). The post hoc analysis revealed significances between shock index >1.5 vs. 0.5 and 0.5 – 1.5 after 24 hours.

The predictive quality of Shock for IL 6 secretion

The predictive values indicative of exsanguination for IL 6 levels in the multiple injured patients were high. The AUC was increasing according to the shock severity (Figure 2). In patients with a high shock index at trauma site with a maximum of 0.975 (Shockindex: >1.5) and in the trauma bay with a maximum of 0.931 (ATLS degree: IV) was reached (Figure 2B) after 48 hours.

The interlink between IL 6 secretion SIRS and Sepsis

The Pearson’s correlation showed a clear correlation between Sepsis and class degree of Shock according to ATLS and for the shock index (Table 2). Independent predictive power of IL 6 in regard to SIRS and sepsis demonstrated significant values only for the IL 6 values after 48 hours (Table 3). The ATLS shock classification revealed independent predictive power for Sepsis, however the shock index did not (Table 3).

Outcome Parameters

Significant differences were found along all groups of the ATLS shock and shock index in this study sample (Table 4). The reduced
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Table 1. Characteristics of the patient sample directly at admission. Dimensions are given in square brackets. AIS: Abbreviated Injury Scale [Range 1-5, arbitrary]. ISS: Injury Severity Scale [Range 16-75, arbitrary]. PT: Prothrombin time, Tc: Platelets. *Kruskal-Wallis H test, †

| ATLS Class | 1 | 2 | 3 | 4 | p-value |
|------------|---|---|---|---|---------|
| Sex [male/female] | 265/90 | 100/43 | 24/6 | 12/5 | <0.001* |
| Age [a] | 40.0 ± 17.8 | 37.5 ± 16.0 | 40.8 ± 16.0 | 34.4 ± 12.8 | 0.389* |
| Time to scene [h] | 6.6 ± 26.1 | 3.4 ± 12.7 | 1.6 ± 0.8 | 2.5 ± 3.7 | 0.597* |
| AIS Head | 3.1 ± 1.8 | 2.7 ± 2.0 | 3.3 ± 1.8 | 3.4 ± 1.6 | 0.235* |
| AIS Face | 0.8 ± 1.2 | 0.6 ± 1.0 | 1.0 ± 1.6 | 0.9 ± 1.3 | 0.122* |
| AIS Thorax | 1.5 ± 1.6 | 2.0 ± 1.7 | 2.2 ± 1.9 | 3.2 ± 1.7 | <0.001* |
| AIS Abdomen | 0.8 ± 1.5 | 1.4 ± 2.0 | 1.8 ± 2.1 | 2.4 ± 1.8 | <0.001* |
| AIS Spine | 0.8 ± 1.4 | 1.0 ± 1.5 | 1.0 ± 1.5 | 1.9 ± 1.8 | 0.005* |
| AIS Extremities | 1.3 ± 1.4 | 1.9 ± 1.6 | 2.0 ± 1.7 | 1.8 ± 1.2 | <0.001* |
| AIS Skin | 0.4 ± 1.0 | 0.8 ± 1.3 | 0.9 ± 1.5 | 1.4 ± 1.7 | <0.001* |
| AIS | 0.4 ± 0.7 | 0.5 ± 0.8 | 0.6 ± 0.8 | 0.8 ± 0.8 | 0.239* |
| ISS | 26.6 ± 11.9 | 32.6 ± 12.3 | 38.1 ± 13.2 | 40.9 ± 13.6 | <0.001* |
| NISS | 36.7 ± 14.5 | 42.1 ± 14.3 | 48.1 ± 12.7 | 50.0 ± 17.5 | <0.001* |
| APACHE II | 13.7 ± 7.1 | 16.0 ± 7.8 | 19.9 ± 8.4 | 19.4 ± 8.5 | <0.001* |
| SAPS II | 30.3 ± 15.1 | 38.3 ± 13.5 | 27.3 ± 12.0 | 31.0 ± 15.5 | 0.344* |
| Base Excess [mM] | -2.6 ± 3.3 | -4.6 ± 4.6 | -6.8 ± 4.7 | -5.7 ± 3.7 | <0.001* |
| Lactate [mM] | 2.5 ± 1.6 | 3.2 ± 2.6 | 4.2 ± 3.2 | 4.2 ± 2.5 | <0.001* |
| Hematocrite [%] | 35.6 ± 7.0 | 32.0 ± 8.5 | 28.0 ± 8.6 | 25.9 ± 8.7 | <0.001* |
| Leukocytes [10^3/µl] | 13.4 ± 5.9 | 13.0 ± 5.9 | 13.3 ± 6.6 | 11.6 ± 6.2 | 0.791* |
| Hemoglobin [g/dL] | 12.2 ± 3.0 | 10.5 ± 3.1 | 9.3 ± 3.0 | 8.8 ± 2.9 | <0.001* |
| Base Excess [mM] | -0.8 ± 1.4 | -1.0 ± 1.5 | -1.0 ± 1.5 | -1.9 ± 1.8 | 0.005* |
| Hematocrite [%] | 35.6 ± 7.0 | 32.0 ± 8.5 | 30.5 ± 8.6 | 28.0 ± 8.6 | <0.001* |
| Leukocytes [10^3/µl] | 80.5 ± 15.4 | 79.0 ± 18.9 | 70.0 ± 20.5 | 62.0 ± 29.5 | <0.001* |
| Tc [10^3/µl] | 117.0 ± 161.0 | 194.6 ± 75.7 | 198.4 ± 110.5 | 146.9 ± 85.0 | 0.024* |

Table 2. Pearson's correlation of IL 6 secretion and of the shock degrees on the development of SIRS and sepsis. IL 6 Levels and shock is clearly correlated with septic complications

| Pearson p-value | SIRS | Sepsis |
|----------------|------|-------|
| IL 6 24h | 0.176 | 0.080 |
| IL 48h | 0.175 | 0.080 |
| ATLS Shock | 0.074 | 0.050 |
| Shock index | -0.107 | 0.090 |

Table 3. Binominal Regression analysis of IL 6 secretion on the development of SIRS and sepsis. IL 6 Level after 48 hours and the ATLS shock degree is clearly associated with septic complication in the multiple injured patient. Hosmer Lemeshow test was > 0.05 for SIRS and sepsis. Data were considered as significant if p < 0.05

| p-value | SIRS | Sepsis |
|---------|------|-------|
| IL 6 24h | 0.958 | 1.000 |
| IL 48h | 0.073 | 1.009 |
| ATLS Shock | 0.093 | 1.362 |
| Shock index | 0.200 | 0.400 |

Table 4. The outcome of the patient sample. The dimensions are given in square brackets. *Kruskal-Wallis H test, †, significance if p < 0.05. Kolmogorov-Smirnov < 0.05 for all groups

| ATLS Class | 1 | 2 | 3 | 4 | p-value |
|------------|---|---|---|---|---------|
| Hosp. [d] | 20.2 ± 13.1 | 28.2 ± 20.3 | 25.6 ± 13.3 | 23.4 ± 20.0 | <0.001* |
| ICU stay [d] | 12.3 ± 10.3 | 16.0 ± 14.3 | 16.3 ± 10.2 | 12.9 ± 11.2 | <0.001* |
| Ventilator [d] | 8.4 ± 9.3 | 10.8 ± 12.7 | 12.0 ± 8.7 | 8.0 ± 6.4 | 0.024* |
| Mortality [%] | 18.3 | 19.1 | 19.2 | 41.7 | 0.508* |

Ventilator days in the ATLS Class IV are probably due to the increased mortality in this patient cohort (Table 4A).

Discussion

SIRS and other multiple trauma-associated complications account for increased hospitalization and worse outcome. In the current study, the question was addressed whether clinical indicators of initial blood loss due to multiple injuries are associated with pro-inflammatory changes. Previous publications in this field discussed the trauma load, the amount of transfusions, plasma expanders and other resuscitative efforts as indicators for the development of SIRS in the multiple injured patient [5,20-22]. From the view of temporal linearity, all pharmacological resuscitative efforts root in the hemorrhage of multiple trauma. The significant results in these studies might be the logical temporal consequence of hemorrhage [5,20-22]. It is hard to distinguish between the blood loss and trauma load as the reason for the pro-inflammatory state. However, even in patients with a higher ISS the blood loss might be negligible as a result self-tamponade and low bleeding as depicted earlier [5,23]. Blood loss in multiple injured patients significantly increased the pro-inflammatory IL 6 level after 24 and 48 hours and was a good predictor for SIRS and sepsis. The pro-inflammatory effect of blood loss as the first event in the timeline after trauma seems to be proven. Many cofounders must be assumed in this study and the transfusion regime as a cofounder cannot be excluded definitively. In 48 hours, a plenty of therapeutic procedures take place in a multiple injured patient. Like the definition of SIRS, transfusions are being applied according to clinical guidelines what makes blood loss and transfusion going hand in hand [24,25]. There are several points of view to blood loss and pro-inflammation. The oxygen supply to all vital systems is reduced leading to an endocrine stress reaction resulting in the secretion of epinephrine and administration of nor-epinephrine by the rescue teams in the hypotonic patient. Maintaining...
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Figure 1. Depicted is the secretion of IL-6 after 24 hours and 48 hours after admission according to the ATLS shock (A) and shock index groups (B). p < 0.05 for IL-6 between the ATLS groups after 24 hours and p < 0.001 between the shock index groups after 24 hours and p < 0.001 for the shock index at trauma site after 48 hours. *Kruskal-Wallis H test. Kolmogorov-Smirnov < 0.05 for all groups. (A and B) Bonferroni p < 0.001, Shockindex >1.5 vs. 0.5-1.5 and <0.5; ATLS shock 4 vs. 3-1

Figure 2. Analysis of the predictive quality of ATLS shock group (A) and shock index group (B) for IL6 secretion if the multiple injured patient. The analysis is given as ROC and given is the area under the curve (AUC), maximal value 1.000

| ATLS Class | 1   | 2   | 3   | 4   |
|------------|-----|-----|-----|-----|
| IL 6 24h AUC | 0.345 | 0.634 | 0.844 | 0.951 |
| IL 6 48h AUC | 0.336 | 0.645 | 0.834 | 0.913 |

| Shock index | <0.5 | 0.5-1.5 | >1.5 |
|-------------|------|---------|------|
| IL 6 24h AUC | 0.443 | 0.557 | 0.963 |
| IL 6 48h AUC | 0.510 | 0.453 | 0.975 |

Conclusions

Class III and IV of Shock and higher levels of shock index states have a higher predictive validity for IL-6 secretion when compared with the shock index at admission in the multiple injured patient. The association between clinical indicators of hemorrhage and IL-6 levels was confirmed, associated with high odds ratio of shock and IL-6 secretion for septic complications. Purposeful clinical consequences lead to quick bleeding control and to damage control transfusion regime.

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Authorship and contributorship

All authors contributed equally to this work in the sense of drafting the manuscript, data collection and revising the draft and being part of the whole intellectual work.

Ethics approval and consent to participate

The study was approved by the Ethics Committee of the University Hospital Zurich with the Permission: "Retrospektive Analysen in der Chirurgischen Intensivmedizin" Nr. St.V. 01-2008.

Conflict of interest

The authors have no conflicts of interest to declare.

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