Histone-DNA Complexes and Coagulation after Intracerebral and Subarachnoid Hemorrhage

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Introduction

Intracerebral hemorrhage (ICH) and subarachnoid hemorrhage (SAH) have limited treatment options and a high mortality. We have previously shown that both conditions are associated with early hypercoagulation.1,2 Histones have been demonstrated to promote thrombin generation.3 In sepsis and trauma patients, studies have revealed an association between histone-DNA complexes and free histone levels and mortality or morbidity.5–6 Theoretically, the effects on coagulation of histones can be investigated in vitro by attempting to block histones with specific antibodies.

The aim of this study was to quantify histone-DNA complex levels in patients with ICH or SAH at the time of admission to hospital compared with 24 hours after symptom onset and to investigate the possible in vitro effect of anti-histone antibodies on thrombin generation.

Materials and Methods

The study was a prospective cohort study. Inclusion criteria were radiologically verified spontaneous ICH or SAH. Patients were excluded if they received medications affecting coagulation prior to inclusion. Further details on the cohort and exclusion criteria have been previously published.1,2 Blood samples were drawn from a peripheral vein or arterial line at admission to hospital and 24 ± 2 hours after symptom onset. Samples were batch analyzed shortly after patient inclusion ended.

Histone-DNA complexes were detected by using the Cell-Death Detection ELISA®PLUS kit (Roche, Mannheim, Germany) and reported in absorbance units (AU). A cohort of 20 healthy men and 20 healthy women established the control group for histone-DNA complexes.7

Thrombin generation was measured in platelet-poor plasma with the addition of tissue factor (5 pM), phospholipids (4 µM), FluCa, hepes, calcium, murine IgG2a kappa isotype control antibody (StemCell Technologies, Vancouver, Canada) and anti-histone antibodies MHIS AB1952 (specific for histone H4) and MHIS AB1992 (specific for histones H1 and H3) kindly donated by Charles T. Esmon, Oklahoma Medical Research Foundation, Oklahoma City, United States. An antibody concentration equivalent to 0.426 mg/mL was chosen for thrombin generation analyses after titration experiments based on available murine studies.8 Four analyses were performed per sample: one with addition of NaCl, one with addition of control antibody and two with the addition of AB1952 and AB1992, respectively. Thrombin generation was quantified with calibrated automated thrombogram (CT, thrombinscope BV, Maastricht, Holland). A cohort of 45 healthy men and 45 healthy women established a control group for thrombin generation results.9

Furthermore, international normalized ratio (INR), activated partial thrombin time (aPTT), antithrombin (functional), thrombin time, fibrinogen (functional, Clauss method), fibrin D-dimer, hemoglobin, platelet and leucocyte count, C-reactive protein (CRP), alanine transaminase, albumin, creatinine, and S100 Calcium Binding Protein B (S100B) were determined.

Descriptive statistics are presented as median with interquartile range (IQR). Paired data were analyzed with a Wilcoxon’s signed rank test (not following Gaussian distribution) or with a Student’s t-test (following Gaussian distribution). Unpaired data were analyzed with a Mann–Whitney test. Correlation tests were performed by using a Spearman’s test. Sample size was defined by the parent study. The primary endpoint was the change in histone-DNA complex levels from admission to 24 hours after symptom onset, for which the power was 100% (mean value 17 AU at admission [standard
deviation 16 AU) and 6 AU at 24 hours) and the significance level, 2α, was 0.05.

The study was conducted in accordance with the Declaration of Helsinki, and the Central Denmark Region Committees on Health Research Ethics approved the study before initiation (case no. 1–10–72–95–14).

**Results and Discussion**

Eighty-seven patients with ICH or SAH were included. The median age was 61 years, and 67% were females. Of the 46 patients with an SAH, 17 underwent surgical coiling and thus received unfractionated heparin. Overall 30-day mortality was 26%, with 33% mortality for SAH patients, and 20% mortality for ICH patients.

Table 1 shows the standard biochemical profiles at admission and 24 hours after symptom onset. Seven patients had a prolonged thrombin time 24 hours after symptom onset due to administration of unfractionated heparin during the coil procedure, close to sample collection. Exclusion of these patients had no significant impact on the results described below.

ICH patients and SAH patients did not differ in histone-DNA complex levels measured at admission ($p = 0.34$) or 24 hours after symptom onset ($p = 0.13$). For the total cohort, histone-DNA complex levels decreased from a median = 11.6 AU (IQR = 6.5–24.5) at admission to median = 3.5 AU (1.5–8.2) 24 hours after symptom onset ($p < 0.0001$). Patients had significantly higher levels of histone-DNA complexes at admission compared with the healthy cohort mean = 5.4 AU (2.0–11.4; $p = 0.0001$).

No difference was demonstrated in histone-DNA complex levels at admission for 30-day survivors (median = 11.3 AU (IQR = 6.4–23.2) compared with nonsurvivors (11.4 AU [5.1–25.9], $p = 1.00$). That histone-DNA complex levels were not found to correlate with mortality is in accordance with a study performed on trauma patients. Thus, we could not confirm the association between levels of histone-DNA complexes and morbidity and mortality found in other patient populations.

<table>
<thead>
<tr>
<th>Parameter</th>
<th>Reference values$^a$</th>
<th>At admission</th>
<th>After 24 hours</th>
<th>$p$-Values</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hemoglobin (mmol/L) ($n_{paired} = 73$)</td>
<td>7.3–9.5</td>
<td>8.4 (7.7–8.9)</td>
<td>7.4 (6.9–8.3)</td>
<td>$&lt;0.0001$</td>
</tr>
<tr>
<td>Platelet count ($\times 10^9$/L) ($n_{paired} = 72$)</td>
<td>165–400</td>
<td>229 (179–268)</td>
<td>212 (170–251)</td>
<td>0.003</td>
</tr>
<tr>
<td>INR ($n_{paired} = 71$)</td>
<td>$&lt;1.2$</td>
<td>1.0 (1.0–1.1)</td>
<td>1.1 (1.1–1.2)</td>
<td>$&lt;0.0001$</td>
</tr>
<tr>
<td>aPTT (s) Before April 5, 2016 ($n_{paired} = 68$)</td>
<td>25–38</td>
<td>28 (26–31)</td>
<td>31 (28–34)</td>
<td>$&lt;0.0001$</td>
</tr>
<tr>
<td>aPTT (s) After April 5, 2016 ($n_{paired} = 9$)</td>
<td>20–29</td>
<td>27 (25–29)</td>
<td>25 (24–31)</td>
<td>0.88</td>
</tr>
<tr>
<td>Antithrombin ($\times 10^3$ IU/L) ($n_{paired} = 37$)</td>
<td>0.80–1.20</td>
<td>0.92 (0.86–1.04)</td>
<td>0.92 (0.85–1.01)</td>
<td>0.84</td>
</tr>
<tr>
<td>Thrombin time (s) ($n_{paired} = 57$)</td>
<td>$&lt;21$</td>
<td>16 (16–17)</td>
<td>16 (15–17)</td>
<td>0.64</td>
</tr>
<tr>
<td>Fibrinogen (µmol/L) ($n_{paired} = 71$)</td>
<td>5.0–12.0</td>
<td>9.2 (7.8–11.1)</td>
<td>9.8 (8.3–10.8)</td>
<td>0.004</td>
</tr>
<tr>
<td>Fibrin d-dimer (mg/L FEU) ($n_{paired} = 71$)</td>
<td>$&lt;0.5$</td>
<td>1.3 (0.66–2.20)</td>
<td>1.2 (0.6–3.5)</td>
<td>0.29</td>
</tr>
<tr>
<td>S100B (µg/L) ($n_{paired} = 71$)</td>
<td>0.02–0.13</td>
<td>0.13 (0.07–0.41)</td>
<td>0.13 (0.06–0.24)</td>
<td>0.33</td>
</tr>
<tr>
<td>CRP (mg/L) ($n_{paired} = 73$)</td>
<td>$&lt;8.0$</td>
<td>2.2 (0.9–4.5)</td>
<td>11.9 (3.9–37.8)</td>
<td>$&lt;0.0001$</td>
</tr>
<tr>
<td>Leucocytes ($\times 10^9$/L) ($n_{values} = 78$)</td>
<td>3.5–10.0</td>
<td>10.9 (7.7–13.4)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Alanine Transaminase U/L ($n_{values} = 84$)</td>
<td>10–70</td>
<td>20 (16–30)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Albumin (g/L) ($n_{values} = 86$)</td>
<td>36–45</td>
<td>38 (35–40)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Creatinine (µmol/L) ($n_{values} = 86$)</td>
<td>45–105</td>
<td>60 (53–75)</td>
<td></td>
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</tr>
</tbody>
</table>

Abbreviations: aPTT, activated partial thrombin time. CRP, C-reactive protein; INR, international normalized ratio; S100B, serum S100 calcium binding protein B.

Note: Values reported as median (interquartile range). As there are missing values (due to failed blood sampling, patient dying before blood sampling etc.), we have reported $n_{paired} = number$ of paired values, $n_{values} = number$ of values. The reference interval for aPTT changed during the inclusion period, and aPTT data are split in two groups accordingly.

$^a$Reference values for adults, combined for men and women, established at the local laboratory.
The main finding of the present study was increased levels of histone-DNA complexes in the acute phase right after an ICH or SAH when compared with 24 hours after symptom onset. A higher thrombin generation was demonstrated in patients at admission than in healthy controls. No convincing effect of adding anti-histone antibodies to thrombin generation could be discerned. Hence, the present study did not support the hypothesis that histones have a significant effect on coagulation as expressed by thrombin generation.

Fig. 1  Thrombin generation with antibody addition in 87 patients with intracerebral or subarachnoid hemorrhage at admission. Thrombin generation in healthy controls and in patients with an intracerebral or subarachnoid hemorrhage, described by lagtime, time to peak, peak and endogenous thrombin potential, with and without added control antibody and anti-histone AB1952 and AB1992 antibodies. Boxes and whiskers indicate median, interquartile range, and range. Number of measurements per group is 81 due to missing values in six patients.
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Conflict of Interest
The authors report no conflicts of interest regarding the present study, but A.M.H. reports other from CSL Behring, Bayer A/S, Boehringer Ingelheim, outside the submitted work.

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References
1 Lauridsen SV, Hvas CL, Sandgaard E, et al. Thromboelastometry shows early hypercoagulation in patients with spontaneous subarachnoid hemorrhage. World Neurosurg 2019;130:e140–e149
10 Johansson PI, Windelev NA, Rasmussen LS, Sørensen AM, Ostrowski SR. Blood levels of histone-complexed DNA fragments are associated with coagulopathy, inflammation and endothelial damage early after trauma. J Emerg Trauma Shock 2013;0(03):171–175