Increased serum levels of interleukin 6 are associated with severe intraventricular haemorrhage in extremely premature infants

A Heep, D Behrendt, P Nitsch, R Fimmers, P Bartmann, J Dembinski

Background: Intraventricular haemorrhage (IVH) and periventricular leucomalacia (PVL) in premature infants presumably have many causes. It has been proposed that inflammatory processes in the fetomaternal unit play an important role in the pathogenesis of these lesions.

Objective: To study the correlation of postpartum serum interleukin 6 (IL6) concentration as a marker of inflammation and neonatal cerebral morbidity in preterm infants < 28 weeks of gestational age.

Methods: A total of 88 infants were grouped according to maximum serum IL6 levels within 12 hours post partum: group A (n = 50), < 100 pg/ml; group B (n = 38), > 100 pg/ml. Ultrasound studies and clinical assessment were performed routinely.

Results: IVH was noted significantly more often in group B (24/38; 63%) than in group A (19/50; 38%) (p = 0.02). In a multiple logistic regression model, raised serum IL6 independently predicted development of severe IVH (odds ratio 8.4; 95% confidence interval 2.85 to 24.9; p = 0.0001).

Conclusions: Raised serum IL6 may serve as a marker for severe IVH in infants < 28 weeks of gestational age. Although cerebral morbidity in premature infants is determined by different variables, the identification of systemic inflammation can help to define the need for anti-inflammatory strategies to prevent cerebral morbidity.

PATIENTS AND METHODS

A retrospective cohort study was carried out on 97 inborn preterm infants, gestational age < 28 weeks, who were admitted consecutively to a single tertiary neonatal intensive care unit at the University of Bonn between January 1999 and December 2001. Nine patients were excluded from analysis because of missing IL6 values (n = 5) or postnatal death within 12 hours (n = 4). The remaining 88 infants were grouped according to maximum IL6 concentration (< 97th centile of healthy neonates) within 12 hours of postnatal age: group A, < 100 pg/ml; group B, > 100 pg/ml. Serum IL6 concentration was measured as part of routine laboratory testing (Immulite; DPC-Biermann, Bad Nauheim, Germany). Clinical and outcome data of the study population were taken from patient records. The clinical risk index for babies (CRIB score) was determined for all study infants from birth weight, gestational age, presence or absence of congenital malformations, worst base excess, and minimum and maximum appropriate fractions of inspired oxygen (FiO2) during the first 12 h of life. All patients were routinely monitored by central or peripheral arterial catheter. Arterial hypotension within the first 12 hours of life was defined as mean arterial blood pressure less than 28 mm Hg. The treatment of arterial hypotension followed a standardised protocol: up to 2 × 10 ml saline infusion/kg body weight/30 minutes, followed by catecholamine treatment (dopamine infusion 5–10 μg/kg/min or adrenaline (epinephrine) infusion 0.1–0.3 μg/kg/min) to maintain blood pressure in the normal range. Cerebral ultrasound was performed with a 8.5–10 MHz transducer (Vingmed Vivid FIVc) at 0–2, 12, 24, 36, 72 hours after birth and at days 7 and 28 of postnatal age. The sonographic findings of IVH and PVL were classified using the criteria given by Volpe.

Abbreviations: IL, interleukin; IVH, intraventricular haemorrhage; PVL, periventricular leucomalacia
parasagittal scan); (c) grade 3 (severe; > 50% of ventricular area in parasagittal scan); (d) apparent periventricular haemorrhagic infarction. According to the classification given by Volpe, in our study severe IVH was defined as either grade 3 IVH or IVH with apparent periventricular haemorrhagic infarction.

Death before postnatal day 28 postnatal and/or severe IVH was defined as a secondary outcome variable.

DATA ANALYSIS
For statistical analysis, we used the Mann-Whitney U test and χ² test with two sided p values to compare values between groups of patients. The effect of serum IL6 > 100 pg/ml at 12 hours of postnatal age and potential confounding variables (gestational age, sex, birth weight, cord blood pH, antenatal steroid exposure, catecholamine treatment, serum C reactive protein concentration, blood leucocyte count) on the incidence of total IVH or severe IVH was tested by stepwise logistic multivariate regression analysis. Data were processed by use of statistical software SPSS 10.7 (SPSS Inc, Chicago, Illinois, USA).

RESULTS
Table 1 summarises the perinatal data of infants in groups A and B. The two groups were similar with respect to gestational age, sex, birth weight, antenatal steroid exposure, arterial umbilical pH, and CRIB score. The variables serum C reactive protein concentration (p = 0.012) and catecholamine treatment (p = 0.002) were measured at 12 hours of postnatal age and were different in the two groups.

In infants with elevated IL6 concentrations (group B), a higher incidence of IVH was noted (p = 0.021). Severe IVH also occurred more often in group B (p = 0.0001). There was a tendency towards a higher rate of survival at day 28 in infants with raised IL6 had a higher risk of death at < 28 days of postnatal age or survival with severe IVH at 28 days of postnatal age (p = 0.0001) than infants with IL6 < 100 pg/ml (group A) (table 2).

In the study population, IVH (43/88) correlated with the variables gestational age (p = 0.003), IL6 > 100 pg/ml (p = 0.034), and catecholamine treatment (p = 0.002) measured at 12 hours of postnatal age. In the first regression model, gestational age (odds ratio (OR) 0.92, 95% confidence interval (CI) 0.86 to 0.98) and catecholamine treatment (OR 3.6, 95% CI 1.15 to 11.3) maintained independent variables associated with the development of IVH.

The second regression model confirmed the independent association of IL6 level > 100 mg/dl and catecholamine treatment with grade 3 IVH or IVH with periventricular white matter disease. The other confounding variables were excluded from the regression model.

DISCUSSION
The results of this clinical study emphasise the relevance of inflammatory processes in the pathogenesis of IVH in extremely premature infants. Infection of the fetomaternal unit increases the risk of preterm delivery and adverse neonatal outcome.5 8 14 28 There is evidence that inflammatory umbilical cord lesions, elevated amniotic fluid, and cord blood IL6 levels are associated with increased neonatal morbidity.5 6 8 11 15 20 IVH and PVL in preterm infants in early postnatal magnetic resonance imaging are apparently related to intraterine T cell activation and increased proinflammatory activity.26 Studies on human cerebral microvascular endothelial cells indicate that these are capable of upregulating inflammatory endothelial mediators in response to proinflammatory cytokines or ischaemia.24 The upregulated expression of adhesion molecules in the endothelial cells, mediating neutrophil rolling and attachment, is the first step of tissue damage. Consecutive pathophysiological steps are vasoparalysis26 and activation of microglial cytokine expression.31–35

Table 2 Outcome data of 88 infants of < 28 weeks gestation

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Group A (n = 50)</th>
<th>Group B (n = 38)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>IVH</td>
<td>19 (38)</td>
<td>24 (63)</td>
<td>0.02</td>
</tr>
<tr>
<td>Severe IVH</td>
<td>6 (12)</td>
<td>21 (55)</td>
<td>0.0001</td>
</tr>
<tr>
<td>PVL</td>
<td>2 (4)</td>
<td>3 (8)</td>
<td>0.64</td>
</tr>
<tr>
<td>Survival day 28</td>
<td>44 (88)</td>
<td>25 (68)</td>
<td>0.077</td>
</tr>
<tr>
<td>Death before day 28 or severe IVH</td>
<td>8 (16)</td>
<td>24 (63)</td>
<td>0.0001</td>
</tr>
<tr>
<td>BPD (FIO2 &gt; 0.21 at 36 weeks gestation)</td>
<td>13 (28)</td>
<td>9 (35)</td>
<td>0.79</td>
</tr>
</tbody>
</table>

Values in parentheses are percentages. Study population divided into two groups according to serum IL6 concentrations at 12 hours of postnatal age: A, < 100 pg/ml; B, > 100 pg/ml. IVH, Intraventricular haemorrhage; PVL, periventricular leucomalacia; BPD, bronchopulmonary dysplasia.

Referring to the development of severe IVH in the study population (27/88; 30%; table 4), we found a correlation with the variables gestational age (p = 0.014), IL6 > 100 pg/ml (p = 0.0001), catecholamine treatment (p = 0.0001), and leucocyte count (p = 0.037).

To determine the influence of inflammation on the development and extent of IVH, stepwise multivariate logistic regression models were applied separately for (a) all patients with IVH (table 3) and (b) patients with severe IVH (table 4).

Table 1 Perinatal data of 88 infants of < 28 weeks gestation

<table>
<thead>
<tr>
<th>Characteristic</th>
<th>Group A (n = 50)</th>
<th>Group B (n = 38)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age (weeks)</td>
<td>24±0</td>
<td>25±1</td>
<td>0.09</td>
</tr>
<tr>
<td>Birth weight (g)</td>
<td>652 (333–1000)</td>
<td>705 (470–910)</td>
<td>0.18</td>
</tr>
<tr>
<td>Male/female</td>
<td>25/25</td>
<td>18/20</td>
<td>1.0</td>
</tr>
<tr>
<td>Antenatal steroids</td>
<td>4/9(%)</td>
<td>39/92(%)</td>
<td>0.43</td>
</tr>
<tr>
<td>Cord blood pH</td>
<td>7.32 (6.9–7.46)</td>
<td>7.32 (7.13–7.44)</td>
<td>0.62</td>
</tr>
<tr>
<td>CRIB score</td>
<td>8(1–21)</td>
<td>8(1–20)</td>
<td>0.063</td>
</tr>
<tr>
<td>Catecholamine treatment</td>
<td>31(62%)</td>
<td>35(92%)</td>
<td>0.002</td>
</tr>
<tr>
<td>C reactive protein</td>
<td>7.14(%)</td>
<td>15.39(%)</td>
<td>0.012</td>
</tr>
<tr>
<td>&gt; 5 mg/l</td>
<td>120–96</td>
<td>1000(114–50000)</td>
<td>0.0001</td>
</tr>
<tr>
<td>IL6 concentration (pg/ml)</td>
<td>34(68%)</td>
<td>32(84%)</td>
<td>0.22</td>
</tr>
<tr>
<td>Mechanical ventilation</td>
<td>48(96%)</td>
<td>38(100%)</td>
<td>0.25</td>
</tr>
</tbody>
</table>

Study population divided into two groups according to serum IL6 level at 12 hours of postnatal age: A, < 100 pg/ml; B, > 100 pg/ml.

*Values are given as median and range.

IL, interleukin; CRIB, clinical risk index for babies; PDA, patent ductus arteriosus.
30 weeks gestation may show a correlation of fetal inflammatory response syndrome (chorioamnionitis, raised pro-inflammatory cord blood cytokines) with reduction of systemic blood pressure, right ventricular cardiac output, and increased incidence of severe IVH. As in the model of cerebral ischaemia-reperfusion injury by Suzuki et al, systemic and local haemodynamic changes and cerebral endothelial damage may contribute to the inflammatory response as postulated in our primary hypothesis.

The study population was defined by gestational age, and not by birth weight, to exclude small for gestational age infants who account for about 20% of our patients with birth weight less than 1000 g. The CRIB score was determined from all study infants to assess the severity of clinical conditions in the first 12 hours of life (fig 1). The median CRIB score did not differ in the two study groups. It is a valid index of initial neonatal risk, predicting neonatal morbidity and mortality even in extremely low birthweight infants as shown by other studies. Statistical analysis of the perinatal data for the two study groups showed no difference in factors known to contribute to the increased risk of neonatal morbidity (antenatal steroids, gestational age, sex, patent ductus arteriosus, mechanical ventilation). Therefore we assume that the elevated serum IL6 concentrations measured at 12 hours of life do not indicate a secondary serum response to perinatal stimuli or represent, irrespective of neonatal inflammation, a sicker infant per se. Owing to the retrospective design of the study, placental histology was not available for all of the study infants to correlate inflammation of the fetomaternal unit with neonatal IL6 serum concentrations and neonatal morbidity.

Studies on the evaluation of IL6 for early diagnosis of neonatal infection indicate that cut off values of 80–100 pg/ml are ideal, detecting early neonatal infection with a high degree of sensitivity and specificity. We chose a cut off value within that range, to exclude other confounding perinatal variables (delivery mode, period of delivery) that might have influenced serum IL6 concentration.

The CRIB score distribution in group A (IL6 < 100 pg/ml) v group B (IL6 > 100 pg/ml) (n = 88) (p = 0.063). The graphic is jittered (5%) to figure overlapping data.
Our data indicate that inflammatory mechanisms may not be separable from other variables that predict cerebral morbidity in a study population of less than 28 weeks gestation. Early neonatal systemic inflammation and haemodynamic disturbances seem to be linked pathophysiological mechanisms which determine the extent of cerebral morbidity in extremely premature infants. Measurement of serum IL-6 concentrations provides important clinical information on early anti-inflammatory processes, before histopathology can confirm fetal involvement in amniotic infection.  

**REFERENCES**