Periventricular leukomalacia in very low birth weight preterm neonates with high risk for neonatal sepsis

Rita C. Silveira,1 Renato S. Procianoy,2 Juliana C. Dill,3 Cristine S. da Costa3

Abstract
Objective: To investigate the association between periventricular leukomalacia (PVL) and neonatal sepsis in very low birth weight infants (VLBW). Methods: We studied VLBW with a clinical suspicion of infection who had been born at our institution between the 1st of August, 2005 and the 31st of July, 2007. Children were excluded if they died before reaching 14 days, had malformations of the central nervous system or congenital infections. Ultrasound brain scans were carried out on the third day and weekly up until the sixth week of life or discharge. Periventricular leukomalacia was diagnosed by persistent diffuse periventricular hyperechogenicity for more than 7 days, or by periventricular cysts. The VLBW were separated into two groups on the basis of the presence or absence of PVL. Sepsis was defined as clinical manifestation plus a positive culture. The Mann-Whitney, chi-square and t tests were applied followed by logistic regression.

Results: A total of 88 VLBW were studied. Of these, 62 (70.5%) survived and 51 (57.8%) had PVL. Both groups were similar in terms of birth weight, gestational age, Apgar score, type of delivery, SNAPPE-II score, presence of necrotizing enterocolitis, persistent ductus arteriosus and deaths. Sepsis and mechanical ventilation were more common in the group with PVL (23.5 and 2.7%, p = 0.005; 86 and 59%, p = 0.004, respectively). Both of these were identified as, independent risk factors for PVL by logistic regression (p = 0.027 and 0.015, respectively).

Conclusions: Chorioamnionitis has been defined as a risk factor for PVL. We have demonstrated that neonatal sepsis is also an important risk factor. We believe that the systemic inflammatory response is the principal factor involved in the etiopathogenesis of PVL among VLBW.

Introduction
The increased survival of progressively more premature newborn infants has resulted in a significant number of preterm infants who grow up with neurocognitive abnormalities.1,2 Currently, periventricular leukomalacia (PVL) is the principal neurological problem affecting children born extremely premature. Approximately 25% of newborn infants with birth weights below 1,500 g who survive to discharge exhibit moderate to severe permanent motor deficits, such as spastic diplegia. At school age, 25 to 50% of children who had been diagnosed with PVL manifest cognitive and learning deficits.3-5

The presentation of PVL is subclinical and so diagnosis is made by neuroimaging examinations (cerebral ultrasound and/or magnetic resonance). An early diagnosis of PVL can be achieved using transfontanelar cerebral ultrasound during the neonatal period used as screening for brain injuries. On the ultrasound brain scan, areas of periventricular hyperechogenicity can be observed which will later develop into periventricular cysts and/or diffuse hyperechogenic areas in the white
matter. This diffuse component of PVL is common among extreme preterm newborns, and the focus appears to be associated with the inflammatory response resulting from ischemia or infection. In these two situations, the factors involved in etiopathogenesis are related to prematurity.

The critical period of cerebral myelination is between 23 and 32 weeks’ postconceptional age, when pre-oligodendrocytes differentiate into immature oligodendrocytes, responsible for initiating myelination of the white matter.

The reduction of cerebral perfusion in very small preterm neonates with deficient cerebrovascular autoregulation is a high risk for the occurrence of injuries to the periventricular white matter. This being so, preterm newborn infants exposed to intrauterine infections are vulnerable to pre-oligodendrocyte death in the face of ischemic insults, even when mild and insufficient to cause injuries by themselves.

Many experimental and clinical studies have demonstrated the link between intrauterine infection and increased occurrence of PVL. A meta-analysis of 26 studies, relating chorioamnionitis to PVL, concluded that, among the newborn infants of mothers with chorioamnionitis, the chances of developing cystic PVL were three times greater. Notwithstanding, studies relating the presence of neonatal sepsis in extreme preterms with increased occurrence of PVL are rare and do not deal with this high risk population specifically.

The objective of our study was to investigate the presence of PVL and the possible risk factors associated with its development, particularly neonatal sepsis, in a population that is especially vulnerable to these conditions: very low birth weight preterm newborn infants (VLBWI).

Methods

This was a prospective cohort study of a population of newborn infants with birth weights between 500 and 1,500 g, born at the obstetrics center at our hospital and admitted to the neonatal intensive care unit (NICU) during the period between 1st August, 2005 and 31st July, 2007. Inclusion was based on the assessment of the duty physician in whose opinion the infants enrolled exhibited clinical signs suggestive of infection during the first week of life.

Exclusion criteria were death before 14 days of life, presence of major congenital malformations or malformation of the central nervous system, congenital STORCH infections (syphilis, toxoplasmosis, rubella, cytomegalovirus or herpes) or HIV+ mother.

An Informed Consent Form was read and signed by parents or guardians before the first cerebral ultrasound scan. The study was approved by Research Ethics Committee at the institution, protocol no. 04/446.

All newborn infants enrolled underwent a first cerebral ultrasound at 3 days of life; and then weekly until the sixth week of life or hospital discharge, if this took place before 6 weeks of life. This screening made it possible to diagnose PVL. Observational clinical monitoring while in NICU allowed for assessment of possible risk factors for PVL. VLBWI were discharged from hospital when they reached a weight of 2,000 g, were able to receive full oral enteral feeding and were clinically stable.

After the serial ultrasound brain scans, patients were divided in two groups:

Group 1: VLBWI with diagnosis of PVL based on diffuse or cerebral cystic white matter injury.

Group 2: VLBWI without diagnosis of PVL, with cerebral echography negative for this condition.

Images were acquired via the anterior fontanelle, with a LOGIQ 500 Scanner (G. E. Medical Systems, United States) and an 11 MHZ transducer. The researchers who carried out the scans were unaware of the patients’ clinical status in terms of presence or absence of sepsis; images were recorded on the machine and reviewed by two researchers who were qualified in cerebral ultrasound scanning.

The criteria for diagnosing PVL were the presence of diffuse periventricular hyperechogenicity that persisted for a period of more than 7 days, without forming cysts (diffuse periventricular white matter injuries) or the presence of cystic lesions of at least 0.5 cm in diameter, distributed bilaterally and located close to the external angles of the lateral ventricles.

The presence of ventricular dilatation without cerebral hemorrhage during any of the serial ultrasound scans was considered to be secondary to necrosis of the white matter present in the diffuse PVL component.

The following data were collected prospectively and later compared between the two groups: sex, birth weight, gestational age, fifth minute Apgar score, SNAPPE-II score, antenatal corticosteroid use, neonatal sepsis, respiratory distress syndrome (RDS), peri-intraventricular hemorrhage (PIVH), non-hemorrhagic periventricular dilatation and pressure positive mechanical ventilation for more than 24 hours. Mortality was also compared between groups.

Gestational age was determined according to the mother’s obstetric history (date of last menstruation) and confirmed by obstetric ultrasound, at 12 weeks’ pregnancy at the latest. In the absence of any reliable maternal data, gestational age was determined by somatic physical and neurological examination of the newborn, using the New Ballard method.

The SNAPPE-II score is a scale to indicate illness severity and risk of mortality during the neonatal period. SNAPP Scores were calculated for all infants within 12 hours of birth.
Infants were diagnosed with RDS if they exhibited expiratory grunting, nostril flaring, sternum retraction, a need for ≥ 40% oxygen and a chest X ray with diffuse ground-glass appearance and a need for exogenous surfactant.

Presence or absence of PIVH was determined from the first ultrasound brain scan and during weekly monitoring scans, according to the classification defined by Papile et al.\textsuperscript{18}

The criteria for diagnosing neonatal sepsis were clinical signs of infection in the presence of microorganism in blood or cerebrospinal fluid cultures. Serum samples for blood culture were obtained using the aseptic collection technique and according to the recommendations laid out by Schelonka et al.\textsuperscript{19}

Clinical signs of infection were the presence of at least three of the findings referred to below, or two clinical findings plus one or more maternal risk factors\textsuperscript{20,21}: thermal instability; apnea, bradyapnea, grunting, tachypnea, sternum and subcostal retraction, nostril flaring and cyanosis; hypotonia and convulsions; irritability and lethargy; gastrointestinal symptoms, such as abdominal distension, vomiting, gastric residues and difficulty accepting food; idiopathic jaundice; cutaneous pallor, cold skin, perspiration, hypotension and capillary refill time longer than 3 s; signs of bleeding, with clinical status suggestive of disseminated intravascular coagulation; subjective assessment: the newborn "does not look well".

Maternal risk factors were obtained from their obstetric records and their medical and perinatal histories. The events most often described were maternal fever, urinary infection and more than 18 hours of rupture of membranes.

The sample size was calculated from published data reporting the occurrence of PVL at 7 to 26% of infants born preterm and weighing less than 1,500 g using proportions. Based on a statistical power of 90% with a level of significance of 5% (α = 0.05) and an estimated PVL incidence of 33% in high risk populations (VLBWI with presumed or confirmed sepsis), the resulting sample size was 30 newborn infants in each group.\textsuperscript{10}

Statistical analysis

Data are presented in the form of mean and standard deviation or median and interquartile range (from the 25th to the 75th percentiles), depending on the behavior of the variable in question. Differences between means were analyzed using Student’s t test, and differences between medians were subject to the Mann-Whitney test. Categorical variables were analyzed using the chi-square test. Variables were included in the logistic regression model if they exhibited statistical significance (p < 0.05) in the comparison between groups, and their odds ratios as risk factors for PVL were calculated. The level of significance was p < 0.05.
Shalak & Perlman have written about the importance of antenatal corticosteroid therapy for reducing severe cerebral hemorrhage, since it accelerates maturation of the germinal matrix, increases systemic arterial pressure with improved cerebral perfusion and appears to be associated with less severe cases of RDS.22 Our data demonstrate that more than half of our VLBWI (58%) had been given antenatal corticosteroid, and that 66% of those did not present hemorrhage or exhibit level I PIVH with later reabsorption, demonstrating the efficacy of this preventative measure for reducing the occurrence of cerebral hemorrhage in premature newborn infants, since those who were given antenatal corticosteroid suffered significantly fewer cerebral hemorrhages.

It has been suggested that antenatal corticosteroid may be protective against injuries to cerebral white matter in VLBWI due to its anti-inflammatory function.23 In our study, a complete course of antenatal corticosteroid was given to 57% of the mothers whose newborn infants developed PVL and to 60% of the mothers of newborn infants without PVL; its administration did not result in a reduced incidence of PVL among VLBWI.

Neonatal sepsis is a clinical condition that activates a cascade of acute inflammatory response mediators. Kadhim et al., in two of their studies, found evidence of significant expression of tumor necrosis factor alpha (TNF-α) and, to a lesser extent, of interleukin-1 beta (IL-1β) in cerebral tissue with PVL, with increased cytokines production in cases where PVL was associated with infection. The inflammatory reaction was detected at an early stage of PVL and extended up to the last phase, cavitation.24,25

### Table 1 - Periventricular leukomalacia in very low birth weight infants and possible risk factors

<table>
<thead>
<tr>
<th>VLBWI (n = 88)</th>
<th>PVL present (n = 51)</th>
<th>PVL absent (n = 37)</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Males (%)</td>
<td>32 (62.7%)</td>
<td>25 (67.5%)</td>
<td>0.44</td>
</tr>
<tr>
<td>Birth weight(g)*</td>
<td>1,010±250</td>
<td>1,068±270</td>
<td>0.29</td>
</tr>
<tr>
<td>Gestational age (weeks)*</td>
<td>29.2±2.5</td>
<td>29.5±2.3</td>
<td>0.15</td>
</tr>
<tr>
<td>Type of delivery</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Vaginal</td>
<td>17 (33.3%)</td>
<td>11 (29.7%)</td>
<td>0.63</td>
</tr>
<tr>
<td>Caesarian</td>
<td>34 (66.7%)</td>
<td>26 (70.3%)</td>
<td>0.63</td>
</tr>
<tr>
<td>Antenatal corticosteroids</td>
<td>29 (57%)</td>
<td>22 (60%)</td>
<td>0.53</td>
</tr>
<tr>
<td>5-minute Apgar score</td>
<td>8 (6-9)</td>
<td>8 (7-9)</td>
<td>0.12</td>
</tr>
<tr>
<td>SNAPPE-II†</td>
<td>18 (7-29)</td>
<td>17 (0-30)</td>
<td>0.097</td>
</tr>
<tr>
<td>Peri-intraventricular hemorrhage (degrees I+II+III+IV)</td>
<td>21 (41%)</td>
<td>13 (35.1%)</td>
<td>0.25</td>
</tr>
<tr>
<td>Ventricular dilatation (n = 15)</td>
<td>12/15 (80%)</td>
<td>3/15 (20%)</td>
<td>0.046</td>
</tr>
<tr>
<td>Neonatal sepsis</td>
<td>12 (23.5%)</td>
<td>1 (2.7%)</td>
<td>0.005</td>
</tr>
<tr>
<td>Respiratory distress syndrome</td>
<td>32 (62.7%)</td>
<td>21 (56.7%)</td>
<td>0.12</td>
</tr>
<tr>
<td>Mechanical ventilation &gt; 24 hours</td>
<td>44 (86%)</td>
<td>22 (59%)</td>
<td>0.004</td>
</tr>
<tr>
<td>Late neonatal mortality</td>
<td>15 (29.4%)</td>
<td>11 (29.7%)</td>
<td>0.45</td>
</tr>
</tbody>
</table>

PVL = periventricular leukomalacia; VLBWI = very low birth weight infants.
Data presented as mean ± standard deviation, median (interquartile range), n (%), chi-square test or Fisher’s exact test when not indicated otherwise.
* Student’s t test.
† Mann-Whitney test.

### Table 2 - Neonatal sepsis and mechanical ventilation as risk factors for periventricular leukomalacia in very low birth weight infants (logistic regression)

<table>
<thead>
<tr>
<th></th>
<th>OR</th>
<th>CI</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>Confirmed sepsis</td>
<td>11.613</td>
<td>(1.42-94.9)</td>
<td>0.027</td>
</tr>
<tr>
<td>Mechanical ventilation</td>
<td>4.3</td>
<td>(1.32-14.43)</td>
<td>0.015</td>
</tr>
</tbody>
</table>

CI = confidence interval; OR = odds ratio.
Prematurity is a considerable neonatal risk factor for PVL, especially for white matter injuries with a diffuse component (non-cystic). There are speculations that other neonatal conditions are associated with diagnosis of PVL, such as perinatal asphyxia, hypovolemia and early neonatal sepsis. Many of these factors cause a reduction in systemic blood pressure, although clinical studies have not demonstrated an association between PVL and postnatal systemic hypotension.22,26,27 It was recently suggested that alterations in cerebral blood flow are a necessary condition for PVL, but are not enough to cause it in isolation. The preference for premature infants is related to the timing of the insult and the distribution of susceptible pre-oligodendrocytes around the brain.8 A question that remains is the influence of the cascade of inflammatory events that occurs in neonatal sepsis. The concern of this study, however, was to exclusively select a population at greater risk of suffering PVL, i.e. sick VLBWI, although not necessarily with a definite diagnosis of neonatal sepsis.

The presence of ischemia, inflammation or systemic infection appears to be etiopathogenic for PVL. It is possible that PVL in preterm newborns is associated with an altered inflammatory response equilibrium within the central nervous system, and that proinflammatory cytokines act directly on the brain, and not as part of a larger systemic response.10 Studies that have assessed the association between maternal chorioamnionitis and PVL have not examined the occurrence of sepsis in premature newborns.11-14 Our study assessed neonatal sepsis as an important risk factor for PVL in VLBWI. We did not assess the presence of chorioamnionitis because we did not have histology for all placentas and, consequently, lacked diagnoses of subclinical chorioamnionitis, which is a risk factor for PVL that has already been described.28

We avoided possible sample selection bias, defining neonatal sepsis as present when newborn infants had a positive culture. For this reason we can safely say that the occurrence of neonatal sepsis in VLBWI increases their chances of PVL by 11 times. In the majority of cases, PVL has a subclinical presentation, and it is therefore of fundamental importance to understand its risk factors, which, together with serial ultrasound brain scans, may make an early diagnosis possible. It is possible to detect PVL by means of transfontanelar cerebral ultrasound, as long as the newborn infant is monitored weekly and up to hospital discharge. This is also useful to predict the development of cerebral palsy.6,29-31 Vollmer et al. studied preterm newborns at 24 to 32 weeks’ gestational age and assessed the efficacy of cerebral ultrasound for detection of cerebral lesions and their relation with delayed neuropsychomotor development at 8 years of age. These authors found that poor prognosis depended fundamentally on the presence and type of lesion found with cerebral ultrasound.26 It is possible that serial cerebral ultrasound scans, in addition to clinical findings from VLBWI in neonatal intensive care units, such as neonatal sepsis, are important prognostic factors.

One possible limitation of our study was that we did not use magnetic resonance imaging to confirm the diagnosis of PVL, especially the diffuse component of white matter injuries. Although magnetic resonance imaging has demonstrated greater diagnostic sensitivity and specificity for PVL, the elevated costs and technical difficulties involved, such as the need for transportation from NICU to radiology department, limits its use as a routine,27 whereas cerebral ultrasound scans are accessible, practical and well-adapted to bedside use.15 We therefore believe that our study makes an additional contribution by demonstrating that it is possible to reach a diagnosis of PVL using ultrasound and following an appropriate follow-up protocol.

We already know that maternal chorioamnionitis is a risk factor for PVL. In this study we demonstrated that neonatal sepsis is also an important risk factor for PVL in VLBWI with clinical suspicion of neonatal infection. It is probable that the systemic inflammatory response to perinatal and neonatal infections is the principal factor involved in the etiopathogenesis of PVL, which is a disease with significant morbidity, in common with cerebral palsy. Further studies attempting to block this inflammatory response may be promising for reducing the magnitude of the PVL and, as a consequence, improve the future quality of life of these VLBWI.

Acknowledgements

The authors would like to thank Vânia Naomi Hirakata and Daniela Benzano, for their help with the statistical analysis.

References


Correspondence:
Renato S. Prociianoys
Rua Silva Jardim, 1155/701
CEP 90450-071 - Porto Alegre, RS - Brazil
Tel.: +55 (51) 3331.5726
Fax: +55 (51) 3331.2738
E-mail: renatosp@terra.com.br