Received Date : 08-Jul-2016
Accepted Date : 02-Aug-2016
Article type : Short Communication

Umbilical Cord CD71+ Erythroid Cells Are Reduced In Spontaneous Preterm Labor

Nardhy Gomez-Lopez,1-3 Roberto Romero,1,4,6 Yi Xu,1,2 Derek Miller,1-3 Ronald Unkel1,2, Tippi MacKenzie7, Michela Frascoli7, Sonia S. Hassan1,2
1Perinatology Research Branch, Program for Perinatal Research and Obstetrics, Division of Intramural Research, Eunice Kennedy Shriver National Institute of Child Health and Human Development, NICHD/NIH/DHHS, Bethesda, Maryland, and Detroit, Michigan, USA
2Department of Obstetrics and Gynecology, Wayne State University School of Medicine, Detroit, Michigan, USA
3Department of Immunology and Microbiology, Wayne State University School of Medicine, Detroit, Michigan, USA
4Department of Obstetrics and Gynecology, University of Michigan, Ann Arbor, Michigan, USA
5Department of Epidemiology and Biostatistics, Michigan State University, East Lansing, Michigan, USA
6Center for Molecular Medicine and Genetics, Wayne State University, Detroit, Michigan, USA
7Eli and Edythe Broad Center of Regeneration Medicine and the Department of

This is the author manuscript accepted for publication and has undergone full peer review but has not been through the copyediting, typesetting, pagination and proofreading process, which may lead to differences between this version and the Version of Record. Please cite this article as doi:10.1111/aji.12556

This article is protected by copyright. All rights reserved
Running head: Umbilical CD71+ erythroid cells in preterm labor

Address correspondence to:

Nardhy Gomez-Lopez, PhD
Department of Obstetrics and Gynecology
Wayne State University School of Medicine
Perinatology Research Branch, NICHD/NIH/DHHS
Detroit, Michigan 48201, USA
Tel (313) 577-8904
Email: nardhy.gomez-lopez@wayne.edu

Perinatology Research Branch, NICHD/NIH/DHHS
Wayne State University/Hutzel Women’s Hospital
3990 John R, Box 4, Detroit, MI 48201, USA
Telephone: (313) 993-2700
Fax: (313) 993-2694
E-mail: romeror@mail.nih.gov

ABSTRACT

This article is protected by copyright. All rights reserved
**Problem:** Preterm neonates are highly susceptible to infection. Neonatal host defense against infection seems to be maintained by the temporal presence of immunosuppressive CD71+ erythroid cells. The aim of this study was to investigate whether umbilical cord CD71+ erythroid cells are reduced in the placenta of women who undergo spontaneous preterm labor/birth.

**Method of Study:** Umbilical cord blood samples (n=155) were collected from the placenta of women who delivered preterm with (n=39) and without (n=12) spontaneous labor, or at term with (n=82) and without (n=22) spontaneous labor. Time-matched maternal peripheral blood samples were also included (n=111). Mononuclear cells were isolated from these samples, and CD71+ erythroid cells were identified and quantified as CD3-CD235a+CD71+ cells by flow cytometry.

**Results:** 1) The proportion of CD71+ erythroid cells was 50-fold higher in cord blood than in maternal blood; 2) a reduced number and frequency of CD71+ erythroid cells were found in cord blood from the placenta of women who underwent spontaneous preterm labor compared to those who delivered preterm without labor; 3) CD71+ erythroid cells were fewer in term pregnancies, regardless of the process of labor, than in preterm gestations without labor; and 4) no differences were seen in umbilical cord CD71+ erythroid cells between women who underwent spontaneous preterm labor and those who delivered at term with labor.

**Conclusion:** Umbilical cord CD71+ erythroid cells are reduced in the placenta of women who undergo spontaneous preterm labor.

**KEYWORDS:** Cord blood, immunosuppression, neonates, parturition, placenta, pregnancy

**INTRODUCTION**

Preterm birth is the leading cause of neonatal morbidity and mortality worldwide, and over 10% of yearly deliveries in the U.S. alone are preterm. Two-thirds of preterm births are preceded by spontaneous preterm labor, a multi-etiological
syndrome causally linked to infection and inflammation. Preterm neonates are at an increased risk for short- and long-term complications. Indeed, preterm neonates are more susceptible to infection than those delivered at term. However, the immune mechanisms underlying this susceptibility are poorly understood.

Neonatal susceptibility was originally attributed to the immaturity of the immune system. However, recent studies are changing this concept. For example, erythroid cells (cells expressing the erythroid-lineage-defining molecule Ter119) are enriched in the neonatal spleen, where they seem to have an immunosuppressive function since they drive Th2 immune responses. Subsequent in vitro and in vivo studies also demonstrated that neonatal CD71+ erythroid cells have immunosuppressive activity in humans (umbilical cord blood) and mice (the neonatal spleen). Such a function is mediated by arginase-2 since supplementation with L-arginine (a substrate for arginase 2) overrides immunosuppression. Neonatal CD71+ erythroid cells seem to participate in the colonization of commensal microorganisms, which occurs shortly after parturition. However, this role is still controversial since CD71 is highly expressed in the gut epithelium. Therefore, the enhanced bacterial clearance that occurs upon anti-CD71 treatment may be the result of diminished intestinal barrier function (i.e. immune priming by leaked microbiota) rather than the absence of CD71+ erythroid cells.

Since CD71+ erythroid cells seem to play a central role in neonatal immunity and premature neonates are highly susceptible to infection, we investigated whether umbilical cord CD71+ erythroid cells were reduced in the placenta of women who underwent spontaneous preterm labor.

MATERIALS AND METHODS

Human subjects, clinical specimens, and definitions

Umbilical cord blood samples were obtained at the Perinatology Research Branch, an intramural program of the Eunice Kennedy Shriver National Institute of Child Health and Human Development, National Institutes of Health, U. S. Department of Health and Human Services (NICHD/NIH/DHHS), Wayne State University (Detroit, MI, USA), and the Detroit Medical Center (Detroit, MI, USA). The collection and utilization of
biological materials for research purposes were approved by the Institutional Review Boards of NICHD and Wayne State University. All participating women provided written informed consent. Umbilical cord samples (n=155) were obtained from the placentas of women who delivered at term with (TIL, n= 82) and without (TNL, n=22) spontaneous labor, or preterm with (PTL, n=39) or without (PTNL, n=12) spontaneous labor. The demographic and clinical characteristics of the four groups of women are shown in Table 1. Umbilical cord blood was collected at birth in ethylene-diamine-tetra-acetic acid (EDTA) containing blood collection tubes by venipuncture of the umbilical vein and then transported to the lab for immediate use. Time-matching maternal blood samples were collected when possible (n=111; TNL = 16, TIL = 60, PTNL = 10, and PTL = 25). Labor was defined by the presence of regular uterine contractions at a frequency of at least two contractions every 10 minutes with cervical changes resulting in delivery. In each case, several tissue sections of the chorioamniotic membranes were evaluated for acute or chronic chorioamnionitis. Also, sections of the umbilical cord were evaluated for phlebitis, arteritis and funisitis. A placental pathology diagnostic was performed according to published criteria by pathologists who had been blinded to the clinical outcomes.

**Cell separation and immunophenotyping**

Mononuclear cells were isolated from blood samples (1 ml) by density gradient using Ficoll-Paque Plus (GE Healthcare Biosciences, Uppsala, Sweden) following the manufacturer’s instructions. Mononuclear cells were collected from the mononuclear layer of the density gradient, washed with 1ml of 1X phosphate-buffered saline (PBS) (Life Technologies, Grand Island, NY, USA) and immediately used for immunophenotyping. Cells were incubated in 100μl of FACS staining buffer (Cat#554656, BD Biosciences, San Jose, CA, USA) for 30 min at room temperature with the following fluorochrome-conjugated anti-human monoclonal antibodies (BD Biosciences): CD3-BV421(clone UCHT1), CD71-APC (clone M-A712) and CD235a-PE (clone GA-R2). Immediately after staining, cells were incubated with 1X FACS Lysing Solution (BD Biosciences) for 2 minutes and washed with FACS staining buffer. Next,
cells were resuspended in 0.5 ml of FACS staining buffer and 10μl of CountBright™ absolute counting beads (Life Technologies) were added. Samples were then acquired using a BD FACS LSR II flow cytometer (BD Biosciences). Data were analyzed using the FACSDiva 6.0 software (BD Biosciences). CD71+ erythroid cells were defined as CD3-CD235a+CD71+ cells\textsuperscript{11}. The total number of CD71+ erythroid cells was calculated from the number of acquired CountBright beads. The flow cytometry figures were prepared using FlowJo software version 10 (TreeStar, Ashland, OR, USA).

**Statistical analysis**

Statistical analyses were performed using SPSS, Version 19.0 (IBM Corporation, Armonk, NY, USA). Normality of the data was tested using the Wilk-Shapiro test. Kruskal-Wallis or Mann-Whitney U- tests were used to compare the medians between groups, and interquartile (IQR) ranges are shown. Comparisons of proportions were made using X\textsuperscript{2} exact tests. A p-value of < 0.05 was used to determine statistical significance. When proportions are displayed, percentages and 95% confidence intervals are shown. Medians are shown with the interquartile range (IQR).

**RESULTS**

A total of 155 umbilical cord blood samples collected from the placenta of women who delivered preterm or at term gestation were included in this study. Demographic and clinical characteristics of the study population are displayed in Table 1. As expected, the gestational age at delivery was shorter in preterm groups when compared to term groups [PTNL 33.7 GWs (IQR = 30.4-35.8) and PTL 35 GWs (IQR = 33.8-36.3) vs. TNL 39 GWs (IQR = 38.8-39.1) and TIL 39.2 GWs (IQR = 38.3-40.3); p<0.001]. Preterm neonates had a lower birth weight than term neonates [PTNL 2032.5 g (IQR = 1260-2520) and PTL 2175 g (IQR = 1767.5-2537.5) vs. TNL 3447.5 g (IQR = 3090-3801.3) and TIL 3172.5 g (IQR = 2882.5-3650); p < 0.001]. All of the term and preterm deliveries without labor were delivered via C-section.

The gating strategy used to determine CD71+ erythroid cells (CD3-CD235a+CD71+ cells) is shown in Figure 1. Briefly, CD71+ erythroid cells were gated within the CD3- gate since T cells (CD3+ cells) also express the CD71 antigen\textsuperscript{17}. CD71+ erythroid cells also expressed the CD235a antigen, the previously used erythroid marker in
umbilical cord blood\textsuperscript{11}. Figure 1 also demonstrates that CD71\(^+\) erythroid cells are abundant in umbilical cord blood; yet, are rare in maternal circulation. There were no differences in the frequency and number of maternal blood CD71\(^+\) erythroid cells among the four groups (data not shown).

The frequency and number of umbilical cord CD71\(^+\) erythroid cells were lower in the placentas of women who underwent spontaneous preterm labor than in those from preterm non-labor controls [PTL 11.10\% CD3-CD235a+CD71\(^+\) cells (IQR = 2.36-21.5) vs. PTNL 37\% CD3-CD235a+CD71\(^+\) cells (IQR = 9.76-43.8), p=0.001; and PTL 50.24 cells/\(\mu l\) (IQR = 8.91-356.15) vs. PTNL 170.81 cells/\(\mu l\) (IQR = 68.88-2759.77), p=0.02; Figure 2]. Neither the frequency nor the number of umbilical cord CD71\(^+\) erythroid cells was different between the placentas of women who underwent spontaneous labor at term and those who delivered at term without labor (Figure 2). No differences were observed in the frequency and number of umbilical cord CD71\(^+\) erythroid cells in the placentas of women who underwent spontaneous preterm labor compared to those who delivered at term with labor (Figure 2).

Umbilical cord CD71\(^+\) erythroid cells were also more abundant in the placentas of women who delivered preterm without labor than in term deliveries with spontaneous labor [PTNL 37\% CD3-CD235a+CD71\(^+\) cells (IQR = 9.76-43.8) and 170.81 cells/\(\mu l\) (IQR = 68.88-2759.77) vs. TIL 8.16\% CD3-CD235a+CD71\(^+\) cells (IQR = 3.19-19.82) and 24.32 cells/\(\mu l\) (IQR = 6.43-97.79); p=0.001 and p=0.0002, respectively; Figure 2] or without spontaneous labor [PTNL 37\% CD3-CD235a+CD71\(^+\) cells (IQR = 9.76-43.8) and 170.81 cells/\(\mu l\) (IQR = 68.88-2759.77) vs. TNL 6.89\% CD3-CD235a+CD71\(^+\) cells (IQR = 3.88-21.42\%) and 29.67 cells/\(\mu l\) (IQR = 5.17-134.89); p=0.01 and p=0.004, respectively; Figure 2].

**DISCUSSION**

Preterm neonates are highly susceptible to infection\textsuperscript{7, 8}. This susceptibility is attributed to the immaturity of multiple immune pathways that are related to arginine depletion\textsuperscript{18}. CD71\(^+\) erythroid cells suppress immune responses via arginase-2 activity\textsuperscript{11}. The study herein demonstrates that CD71\(^+\) erythroid cells are reduced in the umbilical cord of the placentas of women who underwent spontaneous preterm labor compared
to those who delivered preterm in the absence of labor. This data suggests that umbilical cord CD71+ erythroid cells are enriched in preterm gestations, and that the premature process of labor is linked to a reduction in the abundance of these cells. Also, this finding challenges the hypothesis that all preterm neonates have a low frequency of CD71+ erythroid cells, and indicates that the pathological process of labor may reduce immunosuppressive responses in the neonate.

Human CD71+ erythroid cells prevent the activation of other cord blood immune cells in response to heat-killed *Listeria monocytogenes*. The depletion of human CD71+ cells in cord blood samples unleashes the release of TNF-α and IL-6, as well as the activation of CD8+ T cells. Therefore, umbilical cord CD71+ erythroid cells may modulate fetal as well as neonatal immune responses. We found that umbilical cord CD71+ erythroid cells are more abundant in the placenta of women who delivered preterm without labor than in term gestations, regardless of the process of labor. These data suggest that umbilical cord CD71+ erythroid cells are more necessary in fetal than in neonatal immunity. Further research is needed in order to investigate the functional properties of umbilical cord CD71+ erythroid cells in preterm gestations and whether these cells can be detected in earlier stages of pregnancy.

Contrary to the current hypothesis, the study herein demonstrated that umbilical cord CD71+ erythroid cells are not reduced in the placentas of women who underwent spontaneous preterm labor when compared to those who delivered at term gestation with labor. Yet, additional research is needed in order to investigate whether umbilical cord CD71+ erythroid cells display differences in functionality between spontaneous labor at term and preterm labor.

In summary, the study herein demonstrates that umbilical cord CD71+ erythroid cells are reduced in the placentas of women who underwent spontaneous preterm labor when compared to those who delivered preterm without labor. This finding provides insight into the impaired immune mechanisms in premature neonates born from mothers undergoing spontaneous preterm labor. This data also suggests that a reduction in the abundance of umbilical cord CD71+ erythroid cells in preterm gestation could be associated with an increased susceptibility to infection.
ACKNOWLEDGEMENTS
This research was supported, in part, by the Perinatology Research Branch, Division of Intramural Research, *Eunice Kennedy Shriver* National Institute of Child Health and Human Development, National Institutes of Health, U. S. Department of Health and Human Services (NICHD/NIH/DHHS), and, in part, with federal funds from the NICHD/NIH/DHHS under Contract No. HHSN275201300006C. This research was also supported by the Wayne State University Perinatal Initiative in Maternal, Perinatal and Child Health. We thank the physicians and nurses from the Center for Advanced Obstetrical Care and Research, and the research assistants from the PRB Clinical Laboratory, for their help in collecting human samples. We also thank staff members of the PRB Histology and Pathology Units for their examination of the pathological sections, and Tara Mial for her critical readings of the manuscript.

Disclosure/Conflict of Interest
The authors disclose no conflicts of interest.

REFERENCES


**FIGURE LEGENDS**

**Figure 1.** Gating strategy used to identify CD71+ erythroid cells in umbilical cord blood and maternal peripheral blood. CD71+ erythroid cells (CD235a+CD71+ cells) were gated within the CD3- gate.

**Figure 2.** Total frequency and number of CD71+ erythroid cells in umbilical cord blood. Umbilical cord blood were collected from the placentas of women who delivered preterm with (PTL, n=39) and without spontaneous labor (PTNL, n=12), or at term with (TIL, n=82) and without (TNL, n=22) spontaneous labor. Data are shown as scatter plots (median); Mann-Whitney U-tests.
Table 1. Demographics and clinical characteristics of samples used in this study

<table>
<thead>
<tr>
<th></th>
<th>TNL (n=22)</th>
<th>TIL (n=82)</th>
<th>PTNL (n=12)</th>
<th>PTL (n=39)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Maternal age (y; median [IQR]a)</td>
<td>28 (24-29)</td>
<td>24 (21-28)</td>
<td>24 (22-28.3)</td>
<td>23 (20-27.5)</td>
<td>NS</td>
</tr>
<tr>
<td>Body mass index (kg/m²; median [IQR]a)</td>
<td>30 (25.5-35.5)</td>
<td>28 (22-33)</td>
<td>32.1 (27.6-36.9)</td>
<td>27.1 (22.9-33)</td>
<td>NS</td>
</tr>
<tr>
<td>Gestational age at delivery (wk; median [IQR]a)</td>
<td>39 (38.8-39.1)</td>
<td>39.2 (38.3-40.3)</td>
<td>33.7 (30.4-35.8)</td>
<td>35 (33.8-36.3)</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Birth weight (g; median [IQR]a)</td>
<td>3447.5 (3090-3801.3)</td>
<td>3172.5 (2882.5-3650)</td>
<td>2032.5 (1260-2520)</td>
<td>2175 (1767.5-2537.5)</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Baby Sex (n[%])b</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Male</td>
<td>10 (45.5)</td>
<td>38 (46.3)</td>
<td>8 (66.7)</td>
<td>22 (56.4)</td>
<td>NS</td>
</tr>
<tr>
<td>Female</td>
<td>12 (54.5)</td>
<td>44 (53.7)</td>
<td>4 (33.3)</td>
<td>17 (43.6)</td>
<td></td>
</tr>
<tr>
<td>Race (n[%])b</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>NS</td>
</tr>
<tr>
<td>African-American</td>
<td>19 (86.4)</td>
<td>74 (90.2)</td>
<td>8 (66.7)</td>
<td>36 (92.3)</td>
<td></td>
</tr>
<tr>
<td>Caucasian</td>
<td>2 (9.1)</td>
<td>3 (3.7)</td>
<td>3 (25.0)</td>
<td>2 (5.1)</td>
<td></td>
</tr>
<tr>
<td>Hispanic</td>
<td>1 (4.5)</td>
<td>2 (2.4)</td>
<td>0 (0)</td>
<td>1 (2.6)</td>
<td></td>
</tr>
<tr>
<td>Asian</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td></td>
</tr>
<tr>
<td>Other</td>
<td>0 (0)</td>
<td>3 (3.7)</td>
<td>1 (8.3)</td>
<td>0 (0)</td>
<td></td>
</tr>
<tr>
<td>Primiparity (n[%])b</td>
<td>2 (9.1)</td>
<td>15 (18.3)</td>
<td>1 (8.3)</td>
<td>9 (23.1)</td>
<td>NS</td>
</tr>
<tr>
<td>Cesarean section (n[%])b</td>
<td>22 (100)</td>
<td>13 (15.9)</td>
<td>12 (100)</td>
<td>10 (25.6)</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Chronic chorioamnionitis (n[%])b</td>
<td>10 (45.5)</td>
<td>26 (31.7)</td>
<td>5 (41.7)</td>
<td>12 (30.8)</td>
<td>NS</td>
</tr>
<tr>
<td>Acute chorioamnionitis (n[%])b</td>
<td>1 (4.5)</td>
<td>17 (20.7)</td>
<td>0 (0)</td>
<td>8 (20.5)</td>
<td>NS</td>
</tr>
<tr>
<td>Umbilical Cord Pathology (n[%])</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>---------------------------------</td>
<td>--------</td>
<td>--------</td>
<td>--------</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Umbilical Phlebitis</td>
<td>2 (9.1)</td>
<td>20 (24.4)</td>
<td>0 (0)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Umbilical Arteritis</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>5 (12.8)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>Necrotizing Funisitis</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td>0 (0)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*Kruskal-Wallis.*

*χ² test.*

IQR, interquartile range.