Spontaneous Intestinal Perforation is not Associated with the Recent Administration of Antenatal Betamethasone

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Abstract

Intestinal disease in premature infants, especially spontaneous intestinal perforation and necrotizing enterocolitis, contributes significant morbidity and mortality in very low birth weight infants and has a large impact in neonatal care and patient quality of life. In very low birth weight infants, factors influencing a systemic inflammatory response and/ or submucosal thinning, including postnatal glucocorticoids and exposure to indomethacin or ibuprofen, exacerbate the development of intestinal perforation in an already at risk population. In this retrospective analysis we evaluated whether antenatal steroids, a glucocorticoid for fetal lung maturity often given to mothers in close proximity to delivery, may be related to the development of perforation when given close to delivery and without adequate time for recovery of the intestinal mucosa. In our data set, it did not appear to be significantly related. Our results also did not show a categorical association between antenatal steroids and spontaneous intestinal perforation. We did however show a significant relationship between smaller, more depressed infants developing spontaneous intestinal perforation, as well as an association with concomitant sepsis.

Keywords: Spontaneous intestinal perforation; Necrotizing enterocolitis; Antenatal steroids; Betamethasone; Very low birth weight; Fetal infant

Abbreviations: SIP: Spontaneous Intestinal Perforation; NEC: Necrotizing Enterocolitis; ROP: Retinopathy of Prematurity; IVH: Intraventricular Hemorrhage; BPD: Bronchopulmonary Dysplasia; NRBC: Nucleated Red Blood Cell Count

Introduction

Intestinal disease in premature infants, especially spontaneous intestinal perforation and necrotizing enterocolitis, has been an intense focus of current and past research. The significant morbidity and mortality of these diseases creates a large impact in neonatal care and patient quality of life. In very low birth weight infants, factors influencing a systemic inflammatory response and/ or submucosal thinning, including postnatal glucocorticoids and exposure to indomethacin or ibuprofen, exacerbate the development of intestinal perforation in an already at risk population. In this retrospective analysis we hypothesized that antenatal steroids, a glucocorticoid for fetal lung maturity often given to mothers in close proximity to delivery, may be related to the development of perforation when given close to delivery and without adequate time for recovery of the intestinal mucosa (Tables 1 and 2).

Table 1: Comparison by Group.

<table>
<thead>
<tr>
<th>Variable</th>
<th>SIP (N=17)</th>
<th>Control (N=60)</th>
<th>p Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Antenatal Steroids prior to birth (hours)a</td>
<td>32 (3, 240)</td>
<td>51 (1, 696)</td>
<td>NS</td>
</tr>
<tr>
<td>Gestational Age (weeks)b</td>
<td>25.1 (23.2, 28.20)</td>
<td>27.2 (22.0, 35.1)</td>
<td>N=60</td>
</tr>
<tr>
<td>Birth Weight (g)c</td>
<td>697 (300, 1272)</td>
<td>889 (530, 1260)</td>
<td>N=60</td>
</tr>
<tr>
<td>NRBC at birthd</td>
<td>15 (2, 220)</td>
<td>21 (2, 996)</td>
<td>N=60</td>
</tr>
<tr>
<td>Apgar at 5 minutesd</td>
<td>7 (0, 9)</td>
<td>8 (5, 9)</td>
<td>N=60</td>
</tr>
</tbody>
</table>

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several neonatal risk factors have been associated with SIP, little is known about prenatal risk factors. Antenatal indomethacin for tocolysis has been implicated in increasing the risk for periventricular leukomalacia and NEC in premature infants [8]. SIP has also been related to placental inflammation, [9] as well as systemic *Candida* fungal infection. In 2006, Attridge et al. performed a retrospective analysis of 388 infants with SIP and 388 control infants, evaluating for multiple associated variables [10]. They did not find an association between the presence and absence of antenatal steroids.

In this retrospective analysis, we hypothesized that antenatal steroids may be related to the development of perforation when given close to delivery and without adequate time for recovery of the intestinal mucosa.

**Methods**

This study is a retrospective analysis of very low birth weight infants (birth weight ≤ 1500g) who were taken care of at a single institution between August 2006 and December 2011. The study protocol was approved by the institutional review board of Loyola University Medical Center. Our database was queried to identify all infants with SIP. A random sequential number generator was used to identify a cohort of infants ≤ 1500 g without SIP who were born during the same epoch. The medical charts were reviewed for neonatal demographic data, Apgar score at 5 minutes, Nucleated Red Blood Cell Count (NRBC) at birth, blood culture results 48 hours before and after occurrence of SIP, Bronchopulmonary Dysplasia (BPD) defined as oxygen requirement at 36 weeks adjusted gestation, Intraventricular Hemorrhage (IVH), Retinopathy of Prematurity (ROP), as well as the administration of indomethacin, ibuprofen, or postnatal steroids within the first 14 days of life. An infant was considered to have SIP if they had an absence of pneumatosis on radiography, were <3 weeks old, and no surgical or histopathological findings of NEC. Antenatal data was collected on maternal demographics, placental histopathology, and steroid administration prior to delivery. At our institution, betamethasone 12 mg IM every 24 hours is the standard protocol was approved by the institutional review board of Loyola University Medical Center. The study protocol was approved by the institutional review board of Loyola University Medical Center. Our database was queried to identify all infants with SIP. A random sequential number generator was used to identify a cohort of infants ≤ 1500 g without SIP who were born during the same epoch. The medical charts were reviewed for neonatal demographic data, Apgar score at 5 minutes, Nucleated Red Blood Cell Count (NRBC) at birth, blood culture results 48 hours before and after occurrence of SIP, Bronchopulmonary Dysplasia (BPD) defined as oxygen requirement at 36 weeks adjusted gestation, Intraventricular Hemorrhage (IVH), Retinopathy of Prematurity (ROP), as well as the administration of indomethacin, ibuprofen, or postnatal steroids within the first 14 days of life. An infant was considered to have SIP if they had an absence of pneumatosis on radiography, were <3 weeks old, and no surgical or histopathological findings of NEC. Antenatal data was collected on maternal demographics, placental histopathology, and steroid administration prior to delivery. At our institution, betamethasone 12 mg IM every 24 hours is the standard steroid regimen given to mothers with threatened premature birth.

Data were analyzed using SAS version 9.1 (SAS Institute Inc., Cary, NC, USA). Wilcoxon rank sum tests and two sample t-tests were used to examine if there is significant difference in continuous variables between cases and controls. Fisher’s exact tests were used to examine the association between the group and categorical variables.

**Results**

Of the infants at our institution who developed SIP during the time period between August 2006 and December 2011, we identified 17 with a birth weight ≤ 1500 g. A randomized cohort of 60 infants ≤ 1500 g without SIP was also identified. Of the infants with SIP, 33% (18%) did not receive antenatal steroids versus eight (13%) in the control group. There was no significant difference between these two groups using Fisher’s exact tests (p=0.7). There was also no significant difference when comparing the timing of antenatal steroid administration prior to delivery (p=0.25). Mothers of infants who later developed SIP received steroids a median of 32 hours prior to delivery (range 3-240 hours). The control group received steroids approximately 51 hours prior to delivery (range 1-696 hours). Two patients had blood cultures positive for *Candida albicans* in the 48 hours prior to perforation, two had coagulase negative *Staphylococcus* sepsis, and one patient had extended spectrum beta-lactamase positive *Escherichia coli* sepsis. No positive blood cultures were obtained in the control group. There was no placental histopathology available on any patient in the study, no difference in NRBC count was significant between the two groups (median 15 (case) vs 21 (control)).

Table 1 shows the comparison between infants with SIP and control infants. Infants who later developed SIP were significantly more likely to be of younger gestational age (p=0.0004) with a mean gestational age of 25.1 weeks vs. 27.2 weeks in the controls. There was one outlier in the control group with a gestational age of 35.1 weeks.

No difference was significant between genders in the two groups. The case group contained seven females and 10 males while the control group contained 30 females and 30 males. Twins were prevalent in both groups with 41% (n=7) in cases and 27% (n=16) in controls. There was no case of both twins developing SIP. Infants with SIP were also smaller with a mean weight of 697 grams vs. 889 grams (p=0.02), and were more depressed at birth with a 5 minute Apgar median of 7 (range 0-9) vs. 8 (range 3-9) in controls (p=0.04). All infants were <1300 g at birth. No difference was significant between the groups when comparing postnatal steroid administration (p=0.08), although there was a trend towards infants with SIP receiving postnatal steroids earlier in life vs. control infants. Six infants (35%) in the case group received postnatal steroids (five within 48 hours of birth and one 10 days after birth). Of the five infants with very early steroid administration, they received hydrocortisone 3 mg/kg/day. The one infant with steroids on day of life 10 received dexamethasone 0.5 mg/kg/day. Nine patients (15%) in the control group received postnatal steroids in the first one to two weeks of life. More infants with SIP appeared to receive ibuprofen for patent ductus arteriosus than indomethacin vs. the control group (41% (n=7) vs. 18% (n=11)), but we did not see a statistically significant difference between the two groups (p=0.10). Rates of postnatal indomethacin exposure were similar (12% (n=2) vs. 15% (n=9)) (p=1.0). Table 2 denotes outcomes for these infants by time of discharge.

**Discussion**

Intestinal disease in premature infants, especially spontaneous intestinal perforation and necrotizing enterocolitis, has been an intense focus of current and past research. The significant morbidity and mortality of these diseases creates a large impact in neonatal care and patient quality of life. Definitive prevention of SIP and NEC has been elusive. Our understanding of SIP has been growing and many pathophysiological changes have been identified. With SIP appearing to be related to “skewed trophism” and deficiencies of the muscularis propria, [2,4], there exists debate as to whether these changes are acquired, innate, or both. In very low birth weight infants, factors influencing a systemic inflammatory response and/or submucosal thinning, including postnatal glucocorticoids and exposure to indomethacin or ibuprofen, exacerbate the development of intestinal perforation in an already at risk population.

It can be postulated that antenatal steroids, as glucocorticoids, would theoretically contribute to this risk. They have a high rate of administration in the at-risk group and are often given in close proximity to delivery. It has previously been suggested in a large cohort analysis that there is categorically no association between antenatal steroids and SIP. Our hypothesis was that antenatal steroids may be related to the development of perforation when given close to delivery and without adequate time for recovery of the intestinal mucosa. In our small data set, it did not appear to be significantly related. It is possible that a larger case sample would further delineate the relationship, but we presume the effect would be small.

Levels of glucocorticoid bioactivity have been shown to be
elevated >4 fold in newborns that were born <12 hours after the dose of betamethasone was given, a response that resolved if three days passed between steroid dose and delivery [11]. Five of the infants in our case sample received hydrocortisone within 48 hours after delivery, but on further analysis these infants were exposed to antenatal betamethasone between 48-240 hours prior to delivery. Our results also did not show a categorical association between antenatal steroids and SIP, in agreement with Attridge et al. We did however have a significant relationship between smaller, more depressed infants developing SIP, as well as an association with concomitant sepsis. With overwhelming evidence that antenatal steroids reduce the risk of neonatal mortality, RDS, and IVH, the benefits of this medication continue to outweigh the risks [12]. Future research will hopefully further elucidate the pathophysiological triggers for SIP and its avoidance.

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References