Caudal and Penile Blocks Demonstrate Similar Reliability and Efficacy in Pediatric Patients Undergoing Circumcision: A Meta-Analysis

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Abstract

Purpose: Circumcision is one of the most common surgeries performed in the pediatric population. Multiple regional analgesic techniques, including caudal (CB) and penile block (PB), have championed as offering optimal analgesia for circumcision in the post-neonatal pediatric population without clear consensus. This meta-analysis sought to investigate CB and PB’s analgesic efficacy and the impact on postoperative analgesic requirements in pediatric circumcisions. Methods: A comprehensive literature search of PubMed, Google Scholar, and Cochrane Library (1966-2016) was completed to identify all published randomized control trials (RCTs). Keywords searched included “circumcision”, “caudal block”, “penile block”, and “analgesia”. Inclusion criteria were limited to the comparison of PB versus CB in children less than 18 years of age and its efficacy towards circumcision. The efficacy, time to first additive analgesia, time to first micturition, duration of prolonged motor blockade, incidence of vomiting, and length of stay were analyzed. Results: 9 RCTs involving 574 children (N = 287 in CB and PB) were included. No differences in analgesic efficacy (relative risk (RR) = 0.983, 95% confidence interval (CI) = 0.95 to 1.02; p = 0.328) or time to first additive analgesia were observed (standardized difference in mean (SDM) = 0.438, 95% CI = −0.04 to 0.92; p = 0.073). Time to first micturition (SDM = 0.680, 95% CI = 0.40 to 0.96; p < 0.001) and motor block duration (SDM = 0.707, 95% CI = 0.19 to 1.22; p = 0.007) were significantly prolonged in patients receiving CB. No differences were observed between groups in regards to the incidence of vomiting (RR = 1.56, 95% CI = 0.91 to 2.67; p = 0.107) and length of stay (SDM = 0.741, 95% CI = −0.05 to 1.53; p = 0.066). Conclusion: CB and PB offer similar analgesic success rates for pediatric patients (age 18 months to 16 years) undergoing circumcision. CB is associated with a

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trend towards longer duration of analgesia, but is associated with prolonged urinary retention and delayed ambulation. CB use is recommended in non-ambulatory children, whereas PB is recommended in ambulatory children.

Keywords
Circumcision, Caudal Block, Penile Block

1. Introduction
Male circumcision is the most common pediatric surgical procedure performed globally [1] [2]. Worldwide regional prevalence varies significantly due to social, cultural, and religious influences [3]. Circumcision is nearly universal in northern Africa and Muslim Asian countries; whereas prevalence is approximately 15% in sub-Saharan Africa and the United Kingdom [3]. A 2010 report by the Centers for Disease and Prevention (CDC) reported 80% of males between the ages of 14 - 59 years were circumcised in the United States (US) [4] [5]. The CDC estimates that only 58.3% of circumcisions in the US occur during the initial birth hospitalization, resulting in a significant proportion of male children requiring circumcisions as toddlers or adolescents [6].

Reports of medical benefits have contributed to the high prevalence rates of circumcision in the US. Morris et al. (2014) reported that over the course of a circumcised male’s lifetime, the benefits of circumcision exceed the risks by a ratio of 100:1 [5]. The only risks associated with circumcision were surgical complications such as wound infections, whereas the benefits included reduction in urinary tract infections, pyelonephritis, candidiasis, and sexually transmitted infections [5]. The reduced risk of infections is attributed to decreased accumulation of bacteria on the head of penis [5] [7]. Circumcision also reduces the transmission of viruses like human immunodeficiency virus (HIV) and human papilloma virus (HPV) [5]. Additionally, a recent systematic review by Morris and Krieger (2013) involving 19,542 uncircumcised and 20,931 circumcised men demonstrated no adverse effects of circumcision on sexual function, sensitivity, or sexual satisfaction [8]-[10].

Inadequate pain management during circumcision has been associated with altered sensory processing resulting in heightened response to future painful stimuli [11]-[14]. Taddio et al. (1997) reported immunization associated pain increased in circumcised children compared to uncircumcised children [14]. Insufficient analgesia with a placebo during circumcisions was associated with increased future pain sensation in contrast to analgesia with topical anesthetic [14]. Thus adequate analgesia is imperative in circumcision. The most efficient analgesic during circumcision is regional anesthesia, namely caudal (CB) and penile blocks (PB) [15] [16].

CB is a regional anesthesia that is most often used for infra-umbilical incisions, particularly inguinal hernia repair and circumcision [17]. This technique initially uses general anesthesia to sedate the patient and facilitate local anesthetic placement into the caudal epidural space. A study by Shanthanna et al. (2014) reported that landmark ambiguity resulted in up to a 20% technical failure requiring additional postoperative analgesia [18]. Similar to CB, PB is also commonly performed with general anesthesia as it facilitates block placement. There are two types of PB: 1) dorsal nerve penile block which injects a local anesthetic below the pubic bone at the base of the penis and 2) a subcutaneous ring block which injects local anesthetic around the base of the penile shaft [16] [19].

Controversy over the efficacy of these two techniques exists, as some studies have suggested a decreased efficacy and increased block failure rate for CB [18] [20] [21]. A Cochrane systematic review (2008) which included 4 studies concluded that there were no differences in the success rate of analgesic duration between CB and PB, though CB resulted in a longer duration of motor block [22]. This meta-analysis updates the previous Cochrane systematic review (2008) by including five additional randomized control trials (RCT) in an attempt to more precisely define the optimal anesthetic technique for non-neonatal circumcisions.

2. Materials and Methods
2.1. Study Selection
A comprehensive search of all published RCTs comparing CB and PB during circumcision was conducted using PubMed, Google Scholar, and Cochrane Central Registry of Controlled Trials (1966-2015). Additional citations
were searched using references retrieved from prior publications (Figure 1). The last search was conducted on January 28, 2016 and only articles conducted in English were considered. Keywords searched included all combinations of “caudal block”, “penile block”, and “analgesia” in circumcision. The inclusion criteria were limited to RCTs in circumcision, pediatric population (<18 years), comparison of PB and CB, and availability of the event efficacy with sample size. In case of duplicate publications, only the most recent and updated report of the clinical trial was included. The study is compliant with PRISMA guidelines.

2.2. Data Extraction

Articles retrieved from this search were assessed for eligibility and data pertaining to patients, intervention, comparison groups, outcomes, and methodology were abstracted. The primary clinical outcome of interest was efficacy, defined as number of patients requiring no additional pain relief within the first two hours of surgery. Secondary outcomes included time to first additive analgesia, time to first post-circumcision micturition, duration of prolonged motor blockade, risk of vomiting, and length of stay.

2.3. Statistical Analysis

For each trial, relative risk (RR) with a 95% confidence interval (CI) for efficacy and vomiting were calculated. Standard difference in mean (SDM) with 95% CI were calculated for time to first additive analgesia, time to first micturition post-circumcision, and duration of motor blockade. Meta-analysis of the pooled data was performed using the Comparative Meta-Analysis software Version 3 (Biostat, Englewood, NJ). For individual studies reporting zero events in any group, a continuity correction factor of 0.5 was adopted to calculate the RR and variance. In the event of zero events in both groups, the RR was not calculable and the study was excluded from the meta-analysis. Both the fixed effects model and random-effects model were considered, depending on the heterogeneity of the included studies. To assess the heterogeneity between studies, both Cochrane’s Q statistic

Figure 1. CONSORT diagram of the study selection process.
and $I^2$ statistic were used. Heterogeneity was considered statistically significant when $p < 0.05$ or $I^2 > 50$. If heterogeneity was observed, data was analyzed using a random-effects model. In the absence of heterogeneity, a fixed-effects model was assumed.

In the cases of trials reporting results as a median and range, mean and standard deviation was estimated based on Hozo et al. (2005) and Bland (2015) [23] [24]. Outliers as mentioned in the original RCT were not included in the analysis. For all the outcomes, publication bias was first evaluated using a funnel plot, and further evaluated with Egger’s and Begg’s tests. Subgroup analysis was performed based on the anesthetic used, bupivacaine or levobupivacaine. A two-tailed $p$-value of $< 0.05$ was considered statistically significant.

3. Results

3.1. Demographic Characteristics of the Studies

A total of nine RCTs were identified involving 574 children, age 18 months to 16 years (Table 1). The children were equally divided with 287 patients in each group receiving either CB or PB for circumcision. Seven of the nine trials used bupivacaine, while two of the nine trials used levobupivacaine.

Table 1. Characteristics of the randomized control trials comparing caudal block and penile block in circumcision (1966-2015).

<table>
<thead>
<tr>
<th>Author, year</th>
<th>N</th>
<th>Inclusion criteria age range (yrs)</th>
<th>Mean age (yrs) CB group</th>
<th>Mean age (yrs) PB group</th>
<th>Mean weight (kg) CB group</th>
<th>Mean weight (kg) PB group</th>
<th>Medication dose CB group</th>
<th>Medication dose PB group</th>
<th>Time to 1st additive analgesia (min) CB group (mean ± SD)</th>
<th>Time to 1st additive analgesia (min) PB group (mean ± SD)</th>
<th>F/U (hrs)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yeoman et al., 1983</td>
<td>38</td>
<td>1.5 to 12</td>
<td>6.25 ± 3.2</td>
<td>6.5 ± 3.2</td>
<td>22.3</td>
<td>23.2</td>
<td>0.5% bupivacaine (1 mL/yr + 2 mL)</td>
<td>0.5% bupivacaine (1 mL/yr)</td>
<td>-</td>
<td>-</td>
<td>7</td>
</tr>
<tr>
<td>Vater &amp; Wandless, 1985</td>
<td>50</td>
<td>1 to 13</td>
<td>5.6 ± 3.4</td>
<td>4.6 ± 2.9</td>
<td>20.3 ± 7.3</td>
<td>18.1 ± 6.2</td>
<td>0.25% bupivacaine (0.5 mL/kg)</td>
<td>0.5% bupivacaine (1-5 yrs - 3 mL; 6-12 yrs - 4 mL)</td>
<td>313 ± 183.3</td>
<td>256.4 ± 177.6</td>
<td>24</td>
</tr>
<tr>
<td>Irwin &amp; Chang, 1996</td>
<td>50</td>
<td>2 to 12</td>
<td>5.1 ± 2.1</td>
<td>6.4 ± 2.9</td>
<td>18.9 ± 5.9</td>
<td>22.3 ± 9.4</td>
<td>0.25% bupivacaine (0.75 mL/kg)</td>
<td>0.5% bupivacaine (1 mL/kg)</td>
<td>-</td>
<td>-</td>
<td>24</td>
</tr>
<tr>
<td>Mak et al., 2001</td>
<td>126</td>
<td>1 to 12</td>
<td>6.5 ± 2.9</td>
<td>6.5 ± 3.0</td>
<td>24.6 ± 9.4</td>
<td>23.3 ± 8.0</td>
<td>0.25% bupivacaine (0.5 mL/kg)</td>
<td>0.5% bupivacaine (0.5 mL/kg)</td>
<td>354 ± 143.8</td>
<td>366 ± 173.2</td>
<td>96</td>
</tr>
<tr>
<td>Gautlett, 2003</td>
<td>60</td>
<td>1 to 10</td>
<td>5.09 ± 2.32</td>
<td>5.47 ± 2.43</td>
<td>20.9 ± 6.0</td>
<td>22.1 ± 6.9</td>
<td>0.15% bupivacaine (0.5 mL/kg), ketamine (0.5 mg/kg)</td>
<td>0.5% bupivacaine (3 - 5 mL by age + 1 mL subQ)</td>
<td>-</td>
<td>-</td>
<td>24</td>
</tr>
<tr>
<td>Weksler et al., 2005</td>
<td>100</td>
<td>3 to 8</td>
<td>5 ± 2</td>
<td>5 ± 2</td>
<td>20 ± 4</td>
<td>20 ± 4</td>
<td>0.25% bupivacaine (1 mL/kg)</td>
<td>0.5% bupivacaine (0.3 mL/kg)</td>
<td>-</td>
<td>-</td>
<td>24</td>
</tr>
<tr>
<td>Margetts et al., 2008</td>
<td>40</td>
<td>1.5 to 16</td>
<td>7.8 ± 1.7</td>
<td>7.9 ± 1.3</td>
<td>-</td>
<td>-</td>
<td>bupivacaine (0.5 mL/kg), ketamine (0.5 mg/kg)</td>
<td>0.5% bupivacaine (0.25 mL/kg)</td>
<td>485.8 ± 124.9</td>
<td>361.3 ± 183.1</td>
<td>24</td>
</tr>
<tr>
<td>Beyaz, 2011</td>
<td>50</td>
<td>3 to 12</td>
<td>7.4 ± 3.1</td>
<td>8.5 ± 3.5</td>
<td>23.4 ± 8.6</td>
<td>29.4 ±11.3</td>
<td>0.25% levo-bupivacaine (0.5 mL/kg)</td>
<td>0.25% levo-bupivacaine (0.5 mL/kg)</td>
<td>354 ± 15</td>
<td>352 ± 18</td>
<td>6</td>
</tr>
<tr>
<td>Kazak et al., 2012</td>
<td>60</td>
<td>2 to 10</td>
<td>6 ± 3</td>
<td>7 ± 2</td>
<td>23 ± 9</td>
<td>26 ± 6</td>
<td>0.25% levo-bupivacaine (1 mg/kg)</td>
<td>0.25% levo-bupivacaine (1 mg/kg)</td>
<td>458 ± 73</td>
<td>376 ± 68</td>
<td>24</td>
</tr>
</tbody>
</table>

Abbreviations: CB, caudal block; F/U, follow up; hrs, hours; kg, kilograms; mg, milligram; min, minutes; mL, milliliter; N, number of patients included; PB, penile block; SD, standard deviation; subQ, subcutaneous; yrs, years.
3.2. Efficacy of CB and PB in Circumcision

All nine RCTs reported high levels of efficacy for both types of anesthetic block (Figure 2) [15] [16] [20] [21] [25]-[29]. Similar numbers of children had successful blocks with CB and PB (97% and 98% respectively; N = 287 in both arms). Individually, two of the nine trials reported equal efficacy, four studies reported increased efficacy following PB, and three studies reported increased efficacy after CB; however all studies failed to reach statistical significance. There was no significant heterogeneity between trials (p = 0.353, I² = 9.971) and a fixed-effects model was utilized. Meta-analysis revealed no difference in analgesic efficacy between CB and PB (RR = 0.983, 95% CI = 0.95 to 1.02; p = 0.328).

3.3. Time to First Additive Analgesia for CB and PB in Circumcision

Time to first additive analgesia was reported in five trials involving 156 children in the CB group and 157 in the PB group (Figure 3) [15] [16] [21] [26] [29]. Individually, four trials reported longer analgesic duration after CB, two of which reached statistical significance. One of the statistically significant trials used ketamine as an adjuvant to CB [29]. One trial reported equivalent duration of analgesic duration between CB and PB. There was

<table>
<thead>
<tr>
<th>Author/year</th>
<th>Medication</th>
<th>Risk ratio</th>
<th>Lower limit</th>
<th>Upper limit</th>
<th>p-Value</th>
<th>Relative weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>Yeomen,1983</td>
<td>bupivacaine</td>
<td>1.054</td>
<td>0.913</td>
<td>1.216</td>
<td>0.471</td>
<td>5.95</td>
</tr>
<tr>
<td>Vater,1985</td>
<td>bupivacaine</td>
<td>1.043</td>
<td>0.907</td>
<td>1.201</td>
<td>0.553</td>
<td>6.18</td>
</tr>
<tr>
<td>Irwin,1996</td>
<td>bupivacaine</td>
<td>1.085</td>
<td>0.947</td>
<td>1.244</td>
<td>0.241</td>
<td>6.56</td>
</tr>
<tr>
<td>Mak,2001</td>
<td>bupivacaine</td>
<td>0.989</td>
<td>0.918</td>
<td>1.022</td>
<td>0.245</td>
<td>4.31</td>
</tr>
<tr>
<td>Gauntlet,2003</td>
<td>bupivacaine</td>
<td>0.902</td>
<td>0.789</td>
<td>1.030</td>
<td>0.128</td>
<td>4.19</td>
</tr>
<tr>
<td>Welsler,2005</td>
<td>bupivacaine</td>
<td>0.980</td>
<td>0.897</td>
<td>1.028</td>
<td>0.244</td>
<td>26.37</td>
</tr>
<tr>
<td>Beyez,2011</td>
<td>levobupivacaine</td>
<td>1.043</td>
<td>0.907</td>
<td>1.201</td>
<td>0.553</td>
<td>6.18</td>
</tr>
<tr>
<td>Yu,2013</td>
<td>levobupivacaine</td>
<td>0.983</td>
<td>0.949</td>
<td>1.018</td>
<td>0.328</td>
<td>1.37</td>
</tr>
</tbody>
</table>

Abbreviation: CI, confidence interval

Figure 2. Forest plot evaluating the relative risk of caudal and penile block efficacy in circumcision for pediatric patients.

<table>
<thead>
<tr>
<th>Author/year</th>
<th>Medication</th>
<th>Std diff in means</th>
<th>Lower limit</th>
<th>Upper limit</th>
<th>p-Value</th>
<th>Relative weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vater,1985</td>
<td>bupivacaine</td>
<td>0.314</td>
<td>-0.275</td>
<td>0.902</td>
<td>0.296</td>
<td>19.21</td>
</tr>
<tr>
<td>Mak,2001</td>
<td>bupivacaine</td>
<td>-0.075</td>
<td>-0.427</td>
<td>0.277</td>
<td>0.675</td>
<td>23.60</td>
</tr>
<tr>
<td>Margrett,2008</td>
<td>bupivacaine</td>
<td>0.079</td>
<td>0.121</td>
<td>1.460</td>
<td>0.021</td>
<td>17.69</td>
</tr>
<tr>
<td>Beyaz,2011</td>
<td>levobupivacaine</td>
<td>0.121</td>
<td>-0.451</td>
<td>0.693</td>
<td>0.679</td>
<td>19.51</td>
</tr>
<tr>
<td>Kazak,2012</td>
<td>levobupivacine</td>
<td>1.162</td>
<td>0.615</td>
<td>1.710</td>
<td>0.000</td>
<td>19.99</td>
</tr>
<tr>
<td></td>
<td></td>
<td>0.438</td>
<td>-0.040</td>
<td>0.917</td>
<td>0.073</td>
<td>2.00</td>
</tr>
</tbody>
</table>

Abbreviation: CI, confidence interval; diff, difference; Std, standard.

Figure 3. Forest plot evaluating the standardized difference in mean for the time to first additive analgesia with caudal and penile block in pediatric circumcision.
significant heterogeneity between trials \((p = 0.003, I^2 = 75.469)\) and a random-effects model was assumed. Meta-analysis revealed a trend towards prolonged analgesia in CB though results failed to reach statistical significance \((SDM = 0.438, 95\% CI = -0.04 to 0.92; p = 0.073)\).

### 3.4. Time to First Micturition for CB and PB in Circumcision

Time to first micturition was reported in four trials involving 106 patients in each group \(\text{Figure 4} [16] [20] [21] [27]\). All four trials reported longer time to micturition after CB and two trials reached statistical significance. There was no significant heterogeneity between trials \((p = 0.208, I^2 = 34.016)\) and a fixed-effects model was assumed. Meta-analysis revealed longer time to micturition after CB \((SDM = 0.680, 95\% CI = 0.40 to 0.96; p < 0.001)\).

### 3.5. Duration of Motor Blockade for CB and PB in Circumcision

Duration of prolonged motor blockade was reported in four trials involving 99 patients in the CB group and 94 patients in the PB group \(\text{Figure 5} [16] [20] [21] [29]\). Three of the four studies identified a statistically significant prolonged motor blockade after CB, whereas one study reported equivalent duration of motor block in CB and PB. Overall, there was significant heterogeneity between trials \((p = 0.028, I^2 = 67.151)\) and a random-effects model was assumed. Meta-analysis revealed a longer motor block among patients receiving CB \((SDM = 0.707, 95\% CI = 0.19 to 1.22; p = 0.007)\).

### 3.6. Vomiting in CB and PB in Circumcision

Incidence of vomiting was reported in all nine trials but did not include children with failed blocks, resulting in 280 children in the CB group and 283 in the PB group \([15] [16] [20] [21] [25]-[29]\). Two of the nine RCTs reported equal rates of vomiting, three RCTs reported increased vomiting after PB, and four RCTs reported increased vomiting after CB. There was no significant heterogeneity between trials \((p = 0.329, I^2 = 12.7)\) and a fixed-effects model was used. Meta-analysis demonstrated no difference in the RR for vomiting between CB and PB \((RR = 1.56, 95\% CI = 0.91 to 2.67; p = 0.107)\).

### 3.7. Length of Stay Following CB or PB during Circumcision

Length of stay was reported in two of the nine trials involving 80 children in each group \([16] [28]\). Both studies reported longer length of stay after CB, however only one reached statistical significance. There was significant

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**Table 1.**

<table>
<thead>
<tr>
<th>Author, Year</th>
<th>Medication</th>
<th>Std Diff in means</th>
<th>Lower limit</th>
<th>Upper limit</th>
<th>p-value</th>
<th>Relative weight</th>
</tr>
</thead>
<tbody>
<tr>
<td>Vater, 1985</td>
<td>Bupivacaine</td>
<td>0.685</td>
<td>0.096</td>
<td>1.273</td>
<td>0.023</td>
<td>22.41</td>
</tr>
<tr>
<td>Lwin, 1996</td>
<td>Bupivacaine</td>
<td>0.481</td>
<td>-0.093</td>
<td>1.056</td>
<td>0.101</td>
<td>23.52</td>
</tr>
<tr>
<td>Guantlett, 2003</td>
<td>Bupivacaine</td>
<td>1.185</td>
<td>0.621</td>
<td>1.749</td>
<td>0.000</td>
<td>24.43</td>
</tr>
<tr>
<td>Kazak, 2012</td>
<td>Levobupivacaine</td>
<td>0.419</td>
<td>-0.093</td>
<td>0.930</td>
<td>0.109</td>
<td>29.65</td>
</tr>
</tbody>
</table>

**Figure 4.** Forest plot evaluating the standardized difference in mean for time to first micturition with caudal and penile block in circumcision.

**Abbreviation:** CI, confidence interval; diff, difference; Std, standard.

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heterogeneity between trials ($p = 0.017, I^2 = 82.532$) and a random-effects model was assumed. Meta-analysis revealed no difference in mean length of stay between CB and PB (SDM = 0.741, 95% CI = −0.05 to 1.53; $p = 0.066$).

### 3.8. Subgroup Analysis Based on Medication

No difference was observed in regards to bupivacaine or levobupivacaine related efficacy ($p = 0.418$), time to first additive analgesia ($p = 0.528$), time to first micturition ($p = 0.272$), duration of prolonged motor blockade ($p = 0.409$) and risk of vomiting ($p = 0.752$).

### 3.9. Publication Bias

A funnel plot was used to visually assess for publication bias and both Egger’s and Begg’s tests were performed to calculate publication bias (Figure 6). There was no qualitative evidence of asymmetry on the funnel plots. Egger’s and Begg’s tests also revealed no evidence of quantitative publication bias as the lowest $p$-value noted was 0.173 and 0.293, respectively.

### 4. Discussion

Secular circumcision is a common practice in the United States and is associated with decreased sexually transmitted infections and urinary tract infections [5]. The personal medical benefits are reflected in large prevalence of the procedure. In 2012, the Agency for Healthcare Research and Quality reported that hospitalization for circumcision was performed 13.9 times more often than the second most common pediatric surgery, appendectomy [30]. Despite the high prevalence rates of circumcision, perioperative pain management remains a major concern, as a variety of analgesics has rendered the determination of a superior anesthetic difficult [18] [21] [26] [28] [31]-[36].

Analgesic techniques in circumcision include oral sucrose, topical anesthetic, systemic non-steroidal anti-inflammatory drugs (NSAIDs) or opioids, and regional anesthesia [18] [33]-[36]. Non-pharmacological interventions like oral sucrose reduce the duration of cry during circumcision in children less than one year, but appear suboptimal to other anesthetics, as solitary use of oral sucrose is insufficient in treating surgical pain [33]-[37]. Regional anesthetics, in comparison to topical anesthetic and systemic NSAIDs and opioids, offer more optimal pain control with fewer side effects of somnolence, respiratory depression, emesis, and ileus [18] [28] [38]. A one year prospective survey involving 24,409 children reported no long-term complications in children less than
three years and recommended the use of regional anesthesia (overall rate of complication = 0.12%, 95% CI =
0.09 to 0.17) [39]. Despite these advancements, the reliability, effectiveness, and safety of CB to PB in circum-
cision remains controversial [15] [20].

PB provides analgesia over 3/4 of the dorsal penis, while CB provides complete penile analgesia [17] [27] [29]
[31] [40]-[42]. Pain studies have demonstrated longer analgesic duration after CB though this meta-analysis of 9
RCTs (N = 574) failed to demonstrate statistical differences in the efficacy and duration of first additive analg-
esia. These results are consistent with a prior Cochrane systematic review (2008) involving 4 RCTs (N = 336), as
the study also failed to reveal any difference in the efficacy or need for rescue analgesia between CB and PB
(RR = 1.25, 95% CI = 0.64 to 2.44, p = 0.52) [22]. A RCT of 104 patients by Haliloglu et al. (2013) reported a
higher pain score in children postoperatively following PB at 30 minutes but not at 60 minutes (30 minutes: p <
0.001 and at 60 minutes: p = 0.189) [1]. These results suggest any difference in analgesia after PB and CB
wanes quickly after the first half hour of administration.

Analgesic effects in circumcision are varied when bupivacaine and its isomers, levobupivacaine and ropiva-
caine, are compared. Kaya et al. (2012) studied 60 CB patients receiving bupivacaine or levobupivacaine and
found bupivacaine enabled longer analgesic effects compared to levobupivacaine (p < 0.001) [31] [40]. In con-
trast, Locatelli et al. (2005) studied 99 CB patients comparing all three bupivacaine isomers and reported no
difference in analgesic efficacy (p = 0.37) [40]. Subgroup analysis for bupivacaine and levobupivacaine in this
meta-analysis revealed no difference in the measured outcomes, suggesting the use of either isomer is suitable.

CB’s analgesic efficiency can also be improved in concert with adjuvant drugs such as ketamine and magne-
sium. Lonnqvist (2010) reported the use of adjuvant drugs with CB improved analgesic effects up to 24 hours
[17]. Kim et al. (2014) demonstrated time to first additive analgesia, proportion of patients requiring analgesia
postoperatively within 24 hours, and amount of postoperative analgesia were all influenced positively when ad-
juvative drugs were used with CB [42].

In comparison to PB, CB is associated with increased risk of short term complications including urinary re-

![Funnel Plot of Standard Error by Log risk ratio](image-url)
tention and delayed ambulation due to the inhibition of both the sacral parasympathetic and the somatic conduction [20] [21]. Bupivacaine is associated with dose dependent duration of motor blockade [31] [40] [43] [44]. An observational study by Silvani et al. (2006) involving 30 children receiving CB with low concentrations of bupivacaine prolonged analgesia while shortening the duration of motor block (low volume, high concentration analgesic duration: 520 ± 480 min; high volume, low concentration analgesic duration: 952 ± 506 min, p < 0.05) [18] [45]. Optimal anesthetic dose may increase analgesic efficacy after CB while limiting the duration of motor block.

Although the results of this meta-analysis were significant, there are limitations to the study due to the variation and heterogeneity of the RCTs included. The anesthetic type, dose, and adjuvant used varied between studies. The enrollment criteria used in each study differed in regards to age. None of the studies included children less than 18 months, thus reducing the generalizability of these results. The paucity of RCTs in the neonatal population is secondary to the difficulty in pain assessment among newborns, as well as studies associating seizures and arrhythmias to regional anesthesia [17] [39] [46]. That said, more recent studies have documented neonatal complication rates of less than 1/1000 with no long term effects, thereby warranting increased use of CB and PB in this [17] [39] [46].

5. Conclusion

In conclusion, CB and PB regional anesthesia performed during circumcision yielded similar success rates in children aged 18 months to 16 years. CB demonstrated a trend towards longer analgesic duration with significantly more urinary retention and a longer motor block. Results of this meta-analysis suggest CB is a preferred technique compared to PB in non-ambulatory children, as the delayed micturition and ambulation do not significantly impact the length of stay. In ambulatory children, PB should be used over CB to allow for earlier mobility and comfort for the circumcised child. Additional research into bupivacaine isomers and adjuvants should be conducted to determine optimal anesthetic type and dose for pediatric circumcisions, thereby increasing analgesic proficiency while decreasing potential complications.

References


[40] Locatelli, B., Ingelmo, P., Sonzogni, V., et al. (2005) Randomized, Double-Blind, Phase III, Controlled Trial Comparing Levobupivacaine 0.25%, Ropivacaine 0.25% and Bupivacaine 0.25% by the Caudal Route in Children. British Journal of Anaesthesia, 94, 366-371. http://dx.doi.org/10.1093/bja/aei059


