Prednisone for the Treatment of Acute Nonspecific Mesenteric Lymphadenitis

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Abstract

BACKGROUND: Acute nonspecific mesenteric lymphadenitis refers to a common pediatric problem that has no specific treatment.

CASE REPORT: In our study, we describe seven children presenting with abdominal pain as the main symptom of a disease. All patients were treated with prednisone 1 mg/kg (max 40 mg daily) for a maximum of 5 days. In addition, we evaluated the intensity of the pain using a numeric rating scale and achieved a clinically important difference in acute pain relief of 85.3%. After administering corticosteroid therapy, the abdominal pain resolved after 1.7 (1–4) days in all children without any other disturbances.

CONCLUSION: In selective patients with mesenteric lymphadenitis, prednisone can be used as an acceptable form of treatment to reduce the duration of abdominal pain.

Introduction

Primary or acute nonspecific mesenteric lymphadenitis (ANML) is a recognized and well-known cause of acute abdominal pain in children. However, diagnosis of this self-limiting condition is not always straightforward due to the age of the child, lack of specific signs, and common diagnostic criteria [1], [2]. An underlying viral infection (adenovirus, herpes simplex, influenza virus) is thought to be the cause in most cases of ANML. Meanwhile other reported causes of mesenteric lymphadenitis (or secondary) include bacterial (yersiniosis, salmonellosis, and tubercle bacilli), tumor mesenteric lymphadenitis (lymphoma and acute leukemia), parasitic, autoimmune diseases, as well as immunodeficiencies [3]. In clinical practice, these conditions are medical rarities, and the most common dilemma is distinguishing appendiceal inflammation from that of ANML [4]. All children with ANML presented with abdominal pain, whereas other characteristic symptoms include fever, nausea, vomiting, diarrhea, constipation, and anorexia, accompanied with normal or mild elevated markers of inflammatory conditions [4], [5].

In addition to the fact that ANML has no effective therapy, the supportive treatment and time-intensive “wait and see” approach often becomes discouraging for both patients and their families. Against this backdrop, the goal of the current study was to evaluate the efficacy of oral prednisone for pain relief in children with ANML.

Case Report

Seven patients (three boys and four girls) who presented in the Children’s Ambulatory Care Center Subotica between September 2018 and May 2019 with ANML were prospectively analyzed. The average patient age was 8.2 years (range: 5.3–13.7 years). All patients provided a clinical history after which a complete physical examination was performed by pediatrician gastroenterologist and pediatric surgeon. The diagnosis of ANML in children was made in accordance (a) characteristic signs and symptoms; (b) an abdominal ultrasound demonstrating enlarged three or more lymph nodes, short-axis diameter of 8 mm or more in at least one of them (Figure 1); and (c) laboratory and ultrasound exclusion of appendicitis along with other causes of abdominal pain [4], [5].

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The ethical board at our center gave its approval to the study and all parents gave written consent. The patients subjectively evaluated the intensity of the pain using numeric rating scale (NRS) with numbers ranging from 0 to 10 with 0 denoting “no pain” and 10 representing “worst possible pain” as the endpoints [6]. It is notable that we only included patients with pain scores above 6/10 in the study. Pain scores below 4/10 were considered as a satisfactory therapeutic response. Six patients had vomiting (85.7%), five anorexia (71.4%), four fever (57.4%), and one diarrhea (14.2%) and constipation (14.2%). Laboratory tests (complete blood cell count, C-reactive protein-CRP, amylase, liver function tests, urea, creatinine, urinalysis, stool parasite analysis, and stool culture) were found to be normal. During episodes of pain, we recommended acetaminophen for analgesia. Following the exclusion of other diseases, patients were treated with oral prednisone 1 mg/kg (maximum 40 mg) once daily for between 2 and 5 days. The last dose of prednisone was received by patients 1 day after NRS score was found to be below 4/10.

The characteristics of abdominal pain in our children are depicted in Table 1.

All other pre-existing signs and symptoms were found to disappear during the corticosteroid therapy. Even adverse effects of corticosteroids did not show up. Two patients were examined on control ultrasound 1 month after inclusion in the study, whereas mesenterial lymph nodes were normal.

![Image](image.png)

**Figure 1:** Mesenteric lymph node enlargement on abdominal ultrasound in a 7-year-old girl with acute nonspecific mesenteric lymphadenitis

### Table 1: Characteristics of abdominal pain in seven patients affected with acute nonspecific mesenteric lymphadenitis

<table>
<thead>
<tr>
<th>Case number</th>
<th>Localization of abdominal pain</th>
<th>Nocturnal abdominal pain</th>
<th>Duration of pain before treatment (days)</th>
<th>Duration of pain after start of treatment (days)</th>
<th>Pain intensity before treatment (1–10)</th>
<th>Pain intensity after treatment (1–10)</th>
<th>Reduction from baseline pain difference (%)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>Ileocecal</td>
<td>Yes</td>
<td>6</td>
<td>1</td>
<td>9</td>
<td>0</td>
<td>9 (100)</td>
</tr>
<tr>
<td>2</td>
<td>Periumbilical</td>
<td>No</td>
<td>7</td>
<td>2</td>
<td>8</td>
<td>1</td>
<td>7 (87.5)</td>
</tr>
<tr>
<td>3</td>
<td>Periumbilical</td>
<td>No</td>
<td>6</td>
<td>1</td>
<td>8</td>
<td>0</td>
<td>8 (100)</td>
</tr>
<tr>
<td>4</td>
<td>Periumbilical</td>
<td>No</td>
<td>14</td>
<td>4</td>
<td>7</td>
<td>2</td>
<td>5 (71.4)</td>
</tr>
<tr>
<td>5</td>
<td>Periumbilical</td>
<td>Yes</td>
<td>3</td>
<td>1</td>
<td>8</td>
<td>0</td>
<td>8 (100)</td>
</tr>
<tr>
<td>6</td>
<td>Ileocecal and periumbilical</td>
<td>No</td>
<td>3</td>
<td>2</td>
<td>8</td>
<td>2</td>
<td>6 (75)</td>
</tr>
<tr>
<td>7</td>
<td>Ileocecal</td>
<td>Yes</td>
<td>6</td>
<td>1</td>
<td>10</td>
<td>0</td>
<td>10 (100)</td>
</tr>
<tr>
<td>Mean</td>
<td></td>
<td></td>
<td>6.4</td>
<td>1.7</td>
<td>8.2</td>
<td>0.7</td>
<td>7.5 (85.3)</td>
</tr>
</tbody>
</table>

Discussion

As stated before, ANML is a self-limiting disease with mostly no specific cure. The treatment overlaps with many pain syndromes and includes bed rest, hydration (infusion therapy based on the indications), and acetaminophen or a non-steroidal anti-inflammatory agent [7]. To the best of our knowledge, this marks the first attempt to treat ANML with corticosteroids. We searched for the articles published in PubMed, Embase, and Cochrane Library until the month of May 2019 with no language restrictions using the following keywords: “Acute nonspecific mesenteric lymphadenitis,” “mesenteric lymphadenitis,” and “treatment” or “corticosteroids.”

Corticosteroid therapy has been found to be beneficial in some viral infections alone, or as adjuvants with active antimicrobial agents in many infectious diseases [8]. Short-term corticosteroid therapy reduces acute inflammation by inhibiting the activation of inflammatory cells. In addition, it lowers the pain in acute sinusitis as well as upper respiratory tract infections [9]. Studies in adults and children reveal that a single dose of oral dexamethasone in combination with antibiotics increased the likelihood of resolution and improvement of pain in patients with a severe streptococcal sore throat [10]. Correspondingly, Roy et al. found that oral dexamethasone, 0.3 mg/kg for 7 days is effective for pain relief in acute exudative pharyngitis among children with suspected infectious mononucleosis [11]. However, reports pertaining to the effectiveness of corticosteroids for control of swollen lymph glands in mononucleosis are conflicting [12]. Meanwhile corticosteroids do reduce post-operative pain in children and facilitate the prevention of post-operative nausea and vomiting [13].

The controversy is attributed to the potential benefits and adverse events of short-course therapy of corticosteroids in self-limiting diseases such as ANML. At times, health-care providers are reluctant to prescribe corticosteroids due to their immunosuppressive tendencies and potential side effects. However, reports of these side effects during short-course corticosteroid therapy are anecdotal and largely self-limiting (peritonsillar abscess, encephalitis, and myocarditis) [12, 14]. To that end, it is noteworthy that we did not observe any side effects induced by corticosteroid in our patients.
In ANML, children commonly recover within 2–3 weeks, although symptoms persist for 3–10 weeks in half of the patients [7], [15]. Meantime of abdominal pain before therapy in our patients was less than a week; however, the recovery time after treating with prednisone was 1.7 days. We were guided by recommendations of European Society for Paediatric Anaesthesiology that the target should be to achieve and maintain pain scores below the intervention threshold of 4 (on a 10-point scale) [16]. Clinically important differences in acute pain relief using patients’ perception have an impact on which analgesic interventions are regarded useful. The minimum clinically important pain reductions before and after the interventions have varied greatly between studies, with relative difference ranging from 13% to 85% [17]. In our study, the relative difference is 85.3%, which is a large decrease to perceive relief. In addition, we can assume that the resolution of pain and other disturbances can prevent dehydration and obviate the need for infusion therapy.

Notably, we did not use prednisone for all children with ANML in our center, nor do we support their routine use in ANML. We believe that corticosteroid therapy is only needed in severe forms of abdominal pain. Since research on this subject is extremely limited, more placebo-controlled studies with greater number of patients are necessitated to confirm these findings.

Conclusion

Pain control should be a therapeutic goal in patients with ANML. In this regard, prednisone, as adjuvant therapy in conjunction with analgesics, is an adequate and safe therapy, especially in severe forms of this disease.

References