Pediatric pyogenic sacroiliitis and osteomyelitis

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Abstract

Pyogenic sacroiliitis accounts for 1-2% of all cases of septic arthritis with less than 200 cases reported in the English literature since the beginning of the twentieth century. Cultures of joint fluid usually grow Staphylococcus aureus. Prognosis is excellent; however, diagnosis may be difficult due to rarity of disease and non-specific signs, symptoms, and physical findings. Magnetic resonance imaging has been found to be the most useful imaging modality in diagnosis. Most reported cases required prolonged antimicrobial therapy of six to nine weeks. Presented here are two children with pyogenic sacroiliitis managed at a tertiary-care, university hospital and review of the literature on this relatively rare diagnosis.

Case Report #1

A 12-year-old girl presented with a 5-day history of acute onset, non-radiating, right posterolateral hip pain. The pain was initially accompanied by local swelling without erythema and progressively worsened limiting ambulation. There was no history of nausea, vomiting, diarrhea, fever, or rash; however, there was a one-day history of sore throat one week prior to presentation. Past medical, surgical, social, and familial history was non-contributory. Vital signs were notable for a temperature of 38.6 degrees Celsius and heart rate of 123 beats per minute. Physical examination was notable for severe pain at the right hip joint with passive flexion, extension, and internal and external rotation. Neuromuscular function was intact and the remainder of the physical exam was unremarkable. Laboratory studies demonstrated a white blood cell count of 12,200 cells per cubic millimeter with 77.7% neutrophils, anti-streptolysin O antibody of 657 international units per milliliter, erythrocyte sedimentation rate of 84 millimeters per hour, C-reactive protein of 15.1 milligrams per deciliter and a creatinine phosphokinase of 151 international units per liter. Roentogram and ultrasonogram of the right hip were negative. Computed tomography with oral and intravenous contrast of the abdomen, pelvis, and hips was negative. The patient was admitted for further evaluation and begun on intravenous Ceftriaxone, high dose aspirin alone, and morphine. Vancomycin was added when blood cultures became positive for gram positive cocci in clusters (later identified as methicillin-sensitive Staphylococcus aureus) within twenty-four hours. Magnetic resonance imaging of the lumbar spine demonstrated increased joint fluid in the right sacroiliac joint with bone marrow edema in the adjacent right sacrum and increased signal and enhancement of the superior aspect of the right piriformis adjacent to the sacroiliac joint, confirming a diagnosis of septic sacroiliitis associated with piriformis pyomyositis. The patient was evaluated by Pediatric Orthopedic Surgery who determined that surgical debridement was not necessary. The patient was discharged on oral Trimethoprim-Sulfamethoxazole, ambulating with only a slight limp.

The patient was seen in follow-up, approximately one month after discharge and was noted to have persistent right hip pain requiring a walker for ambulation. The erythrocyte sedimentation rate had increased to 70 millimeters per hour from 44 at discharge. A repeat magnetic resonance imaging (MRI) and computed tomography (CT) showed no evidence of a sequestrum. Poor patient compliance was noted due to nausea; antibiotics were changed to Cephalexin and Rifampin. One month later the patient experienced a significant decrease in pain and was ambulating without an aid. Erythrocyte sedimentation rate (ESR) was 17 and antibiotics were continued for a total course of three months with full recovery.

Case Report #2

A previously healthy 23-month-old male presented with four separate episodes, over a 2 month period, of refusing to bear weight on his right lower extremity. The first episode lasted for eight hours, self-resolved, but recurred a couple of weeks later. Each episode had a sudden onset, lasted a few hours longer than the previous episode, and then resolved. There was no fever or other associated symptoms or physical finding. There was no history of trauma or recent recent illnesses.

Bilateral roentograms of lower extremities during the second episode were normal. An MRI of his right lower extremity obtained during the third episode was normal. However, a CT of the right sacrum demonstrated sclerosis and periosteal irregularity of the right superior or sacroiliac joint. An MRI of the right sacrum revealed a focus of low T1 and high T2 signal in the upper aspect of the right sacrum, and the patient was referred to pediatric orthopedic surgery and infectious diseases.

Discussion

The evaluation of a child with a limp involves consideration of several potential etiologies including infection, fracture, and rheumatic arthritis. Septic sacroiliitis, also known...
as pyogenic sacroiliitis (PS), is an uncommon diagnosis in the pediatric patient, accounting for one to two % of all cases of septic arthritis, with fewer than 200 cases of PS having been reported in the English literature since the beginning of the twentieth century.

Schaad et al. reported 77 patients with sacroiliitis between 1961 and 1980 noting that infection was more common in boys during late childhood and in boys often presenting sub-acutely. Bone scans were generally helpful in making the diagnosis, and aspiration of the joints frequently revealed *Staphylococcus aureus* as the infecting agent. Management frequently required antibiotic targeted towards *Staphylococcus aureus* and surgical debridement or drainage and the prognosis was excellent.

Since 1980, there have been six case series which included five or more children (<18 years old) that include outcomes. *Staphylococcus aureus* was the most frequently recovered pathogen (12 of 16; 75% including our cases); however, case reports have also identified *Salmonella* and *Brucella* as less frequent organisms. Previously there were a total number of 39 patients in the six case series (Table 1) most of whom were older than 10 years, although the youngest reported child was 20 months old. Preceding trauma (9 of 25)

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**Table 1. Patient data.**

<table>
<thead>
<tr>
<th>Article</th>
<th>Number of children</th>
<th>Age range (years)</th>
<th>Male:Female</th>
<th>Preceding trauma</th>
<th>Preceding infection</th>
<th>Duration of Sx</th>
<th>WBC range</th>
<th>ESR range</th>
<th>Blood culture</th>
<th>Aspirate culture</th>
<th>Initial X-ray</th>
<th>Initial CT</th>
<th>Initial Bone scan</th>
<th>MRI</th>
<th>Abscess</th>
<th>Antibiotic</th>
<th>Outcome</th>
<th>Residual X-ray changes</th>
</tr>
</thead>
<tbody>
<tr>
<td>Schaad</td>
<td>6</td>
<td>2-12</td>
<td>4:2</td>
<td>1/6</td>
<td>NR</td>
<td>1.5wks-6wks</td>
<td>6.1-14</td>
<td>50-127</td>
<td>NR</td>
<td>Pos /4/ not specified</td>
<td>2/5</td>
<td>ND</td>
<td>4/5</td>
<td>ND</td>
<td>0/6</td>
<td>Parenteral Flucloxacillin and Fucidin for 6 weeks</td>
<td>Cured</td>
<td>0/6</td>
</tr>
<tr>
<td>Ncube</td>
<td>6</td>
<td>6-14</td>
<td>1:5</td>
<td>3/6</td>
<td>2/3</td>
<td>2d-7d</td>
<td>9.0-17.0</td>
<td>31-98</td>
<td>3/6 positive for <em>Staphylococcus aureus</em></td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>ND</td>
<td>ND</td>
<td>ND</td>
<td>0%</td>
<td>0/6</td>
<td>Cured</td>
<td>0%</td>
</tr>
<tr>
<td>Moyer</td>
<td>6</td>
<td>1.5-15</td>
<td>5:1</td>
<td>3/6</td>
<td>3/6</td>
<td>9d-28d</td>
<td>6.6-16.6</td>
<td>55-106</td>
<td>3/6 positive for <em>Staphylococcus aureus</em></td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>0%</td>
<td>4/5</td>
<td>4/5</td>
<td>0/6</td>
<td>0/6</td>
<td>Cured</td>
<td>0/6</td>
</tr>
<tr>
<td>Abbott</td>
<td>9</td>
<td>20mo-14</td>
<td>5:4</td>
<td>NR</td>
<td>4/9</td>
<td>4d-28d</td>
<td>6.0-22.1</td>
<td>10-110</td>
<td>3/6 positive for <em>Staphylococcus aureus</em></td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>0%</td>
<td>ND</td>
<td>ND</td>
<td>0%</td>
<td>0/6</td>
<td>Cured</td>
<td>0%</td>
</tr>
<tr>
<td>Carly</td>
<td>7</td>
<td>2-16</td>
<td>2:5</td>
<td>2/7</td>
<td>2/7</td>
<td>4-28d</td>
<td>8.9-16.7</td>
<td>49-88</td>
<td>4/7 positive for <em>Staphylococcus aureus</em></td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>0%</td>
<td>ND</td>
<td>ND</td>
<td>0%</td>
<td>0/6</td>
<td>Cured</td>
<td>0%</td>
</tr>
<tr>
<td>Aprin</td>
<td>10</td>
<td>12-15</td>
<td>2:5</td>
<td>1/7</td>
<td>1/7</td>
<td>4d-28d</td>
<td>5.2-14.1</td>
<td>40-88</td>
<td>2/3 positive for <em>Staphylococcus aureus</em></td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>0%</td>
<td>ND</td>
<td>ND</td>
<td>0%</td>
<td>0/6</td>
<td>Cured</td>
<td>0%</td>
</tr>
<tr>
<td>Turen</td>
<td>11</td>
<td>20mo-16</td>
<td>2:3</td>
<td>19/20</td>
<td>1/1</td>
<td>12-19</td>
<td>5.2-22.1</td>
<td>12-16</td>
<td>2/2 (one low back pain)</td>
<td>Negative Blood culture</td>
<td>0%</td>
<td>0%</td>
<td>ND</td>
<td>ND</td>
<td>0%</td>
<td>0/6</td>
<td>Cured</td>
<td>0%</td>
</tr>
<tr>
<td>Subtotal</td>
<td>41</td>
<td></td>
<td></td>
<td></td>
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</table>

ND, not done; NR, not reported.
or infection (9 of 28) was previously reported in about one-third of the children previously reported. Neither of our patients had a preceding infection or traumatic injury. Patients presented both acutely or after a subacute course. As with our patients, signs and symptoms were relatively nonspecific, and patients were often admitted with alternative diagnoses. The physical exam was often nonspecific, with difficulty in localizing pain to the SI joint. Aprin and Turen\textsuperscript{11} noted that 7 of 7 patients were positive in the FABERE test. This involves hip flexion, abduction, external rotation, and extension, placing the lateral malleolus on the contralateral knee. Pressing down gently on the ipsilateral knee in this position, stresses an inflamed SI joint and causes pain. Similarly, Ncube\textsuperscript{4} noted that five of six patients had a positive sacroiliac stress test.

Laboratory examinations are non-diagnostic in pyogenic sacroilitis tending to be variable. White blood cell count and ESR were sometimes elevated but not consistently. Blood cultures were positive in 16 of 32 (50%; including our 2), while SI joint aspirate cultures were positive in 12 of 15 patients, (67%; including our 2). In the six case series, as well as with our two patients, \textit{Staphylococcus} species was the most frequently isolated organism. Coagulate negative \textit{Staphylococcus} is far less pathogenic than \textit{S. aureus} and may be a contaminant; however it is frequently isolated from patients with chronic osteomyelitis\textsuperscript{12} and may explain the slow progression of symptoms in case #2. \textit{Escherichia coli} were identified in two cases and \textit{Streptococcus pneumonia} in one case.

Roentograms of the hips are generally non-diagnostic in pyogenic sacroilitis (3 of 30 including current cases). CT scans were positive in 12 of 15 (including current cases) and bone scans in 28 of 35 cases. The study by Doita et al.\textsuperscript{13} is the only series to include MRI findings which were positive in 5 of 5 cases (7 of 7 including current cases). The authors noted that both MRI and bone scan were sensitive for PS; however, bone scans may be negative if done too early after onset of symptoms. Indeed, Moyer et al.\textsuperscript{5} noted one case where an equivocal bone scan became positive when repeated 30 days later and another where a negative bone scan became positive four days later. In addition, bone scan cannot differentiate PS from psosas or gluteal abscesses or identify infection of adjacent tissue; therefore, MRI may provide more diagnostic accuracy, achieving high sensitivity and specificity.\textsuperscript{13}

Traditionally extended-spectrum penicillins or first generation cephalosporins have been the antibiotic of choice. However, methicillin-resistant \textit{Staphylococcus aureus} will require Clindamycin, Linazolid, or Trimethoprim-Sulfamethoxazole. Most patients had excellent outcomes, although, residual asymptomatic roentogram changes were noted in 12 of 24 (including current cases).

Abscesses were reported in 3 of 30 (including current cases) patients. Though there is little pediatric data with respect to surgical drainage, Hodgson\textsuperscript{14} reported a series of 12 adult patients, of whom 11 had surgical drainage. All had a good outcome, though several had mild residual symptoms and one developed a draining sinus, in contrast to a 30-40% mortality rate prior to the antibiotic era.\textsuperscript{2}

### Conclusions

Pyogenic sacroilitis is a relatively uncommon diagnosis in pediatrics. Signs and symptoms can be nonspecific, and physical examination is often unhelpful. The FABERE test is thought to be useful in evaluating the sacroiliac joint. When inflammatory parameters are elevated they are consistent with the diagnosis, but their absence is not reassuring. Blood cultures and aspiration of the SI joint are useful in identifying the infecting agent. While imaging may include roentogram, bone scan, CT, and MRI with varying degrees of sensitivities, the MRI may be the most sensitive in securing the correct diagnosis. Antibiotic therapy initially should be started empirically towards \textit{Staphylococcus aureus} and then continued for at least three weeks. The optimum duration of therapy is not known, though in most cases antibiotics were continued for six to nine weeks.

In the first case presented, noncompliance with Trimethoprim-Sulfamethoxazole became an issue resulting in a prolonged clinical course requiring a longer total duration of antibiotic therapy. The second case presented subacutely, taking two months to establish a diagnosis. It is unclear whether the patient initially had sacroilitis that progressed into bony involvement of the sacrum or whether the sacrum was involved first and that progressed to involve the sacroiliac joint. Because of the chronic osteomyelitis, antibiotic treatment was extended in this patient. Consistent with patients previously described in other reports both patients ultimately responded to therapy with good final outcomes.

### References