1.0 Introduction

Acute osteomyelitis (OM) is most often a consequence of hematogenous spread of bacteria from the skin or mucosal surfaces. The most common causative pathogens in otherwise healthy children are Staphylococcus aureus and Kingella kingae, the latter primarily impacting children 6 months to 4 years of age. Other pathogens include Streptococcus pneumoniae and Streptococcus pyogenes. While OM can occur in any bone, it is more common in the metaphysis of long tubular bones. The predominant symptom is pain, with children presenting with pseudoparalysis (decreased movement) of the affected limb or limping. Fever may also accompany OM but is not necessary for diagnosis.

The gold standard for diagnosis of OM is pathological assessment of a bone specimen. However, in practice this test is invasive and usually not performed. Magnetic resonance imaging (MRI) is a sensitive non-invasive technique often used to confirm the diagnosis of OM. Laboratory markers, such as white blood cell (WBC) count and C-reactive protein (CRP) can be used in conjunction with clinical features and imaging to support the diagnosis.

All patients with OM require admission to hospital for IV antibiotics and evaluation by a multidisciplinary team, usually consisting of Paediatric Medicine, Infectious Diseases and Orthopaedics. Patients with uncomplicated OM are usually started on IV cefazolin, which provides good coverage for methicillin susceptible S. aureus and K. kingae. In children older than 4 years of age where K. kingae coverage is less important, cloxacillin or cefazolin are reasonable options for empiric therapy. Once the child’s clinical status has improved and inflammatory markers are improving, children are transitioned to oral antibiotics to complete the treatment course.

The purpose of this Clinical Practice Guideline is to standardize care for patients with acute osteomyelitis, facilitate collaboration between Paediatric Medicine, Infectious Diseases and Orthopaedics, and improve outcomes.

2.0 Definitions

- Acute osteomyelitis: Infection of the bone with symptom duration < 2 weeks.
- Subacute osteomyelitis: Infection of the bone with symptom duration > 2 weeks
- Chronic osteomyelitis: Infection of the bone with symptom duration > 1 month.
• Septic arthritis: Infection of the joint synovial fluid.

Target Users

The target users of this Clinical Practice Guideline include, but are not limited to:

• Paediatric Medicine providers
• Infectious Diseases providers
• Orthopaedic Surgery providers
• Plastic Surgery providers
• Emergency Medicine providers
• Pharmacists
• Radiology and Image Guided Therapy

Target Population

Inclusion criteria: This inpatient management pathway is primarily intended for use in clinically stable children ≥3 months of age with a diagnosis or suspected diagnosis of acute OM. For initial management in the ED, see: ED Pathway for Suspected Acute Osteoarticular Infection

Exclusion criteria: The Clinical Practice Guideline is not intended for use in patients who:

• Are < 3 months of age
• Have sickle cell disease (inclusive of any genotype)
• Are immunocompromised (including asplenic patients)
• Are hemodynamically unstable (e.g., septic shock)
• Have chronic or subacute osteomyelitis
• Have a diagnosis of osteomyelitis of the frontal bone (Pott puffy tumor), spine, jaw, or skull
• Have a diagnosis of acute multifocal osteomyelitis or recurrent osteomyelitis
• Have a diagnosis of osteomyelitis involving the digits or other bones of the hand. These infections are not usually acute, and most often present with pain and swelling without fever. Plastic surgery should be engaged for hand infections
3.0 Guideline

**Osteomyelitis Pathway**

Admission to Paediatric Medicine for possible osteomyelitis

- Start high dose IV cefazolin (refer to e-Formulary) and order MRI
  - Ensure blood culture drawn prior to initiating antibiotics
  - MRI should occur within 24h
  - Refer to e-Formulary for empiric antibiotic guidelines if MRSA risk factors

MRI confirms osteomyelitis

Yes

Daytime consult to Infectious Diseases and Orthopaedics¹
  - Consult should be completed within 24-48h
  - ID to provide additional antibiotic recommendations as needed and organize ID clinic follow up
  - Orthopaedics to determine whether surgical intervention is indicated
  - Plastics Surgery should be consulted for any hand osteomyelitis

Blood culture negative

Yes

Continue on IV antibiotics until discharge criteria met (see Table 1).

No

Blood culture growing MSSA
  - Continue on IV antibiotics until discharge criteria met (see Table 1).
  - Repeat blood cultures and if repeat cultures are negative and patient is improving with an otherwise uncomplicated osteomyelitis, patient can be transitioned to oral antibiotics as per Table 2 when discharge criteria met (patient does not require 14 days IV antibiotics as in MSSA sepsis without a source).
  - If there are any complicating factors (e.g. abscess, poor response to antibiotics) the primary team should discuss with Infectious Diseases regarding oral stepdown

Blood culture growing a bacteria other than MSSA
  - Discuss with Infectious Diseases

Does patient meet discharge criteria, as per Table 1?

Yes

Step down to oral antibiotics as per Table 2
  - Repeat CRP once prior to discharge to ensure it is trending down
  - Discharge home with follow up with Infectious Diseases as an outpatient +/- Orthopaedics as needed
  - If hand osteomyelitis then Plastics Surgery will arrange follow up

No

Consider alternate etiology

No

Continue IV antibiotics and reassess in 24-48h
Table 1: Discharge Criteria
If discharge criteria is met, switch to oral antibiotics as per Table 2

- Clinical improvement
  - Afebrile for > 24hrs
  - Significant reduction in pain and local signs of inflammation
  - Improved musculoskeletal function (e.g., improved range of motion of affected area; with lower extremity there should be an ability to weight bear, with upper extremity infection there should be only mild pain with routine use) 1, 2
- AND down trending CRP

1. Clinical Practice Guideline by the Pediatric Infectious Diseases Society and the Infectious Diseases Society of America: 2021 Guideline on Diagnosis and Management of Acute Hematogenous Osteomyelitis in Pediatrics 2021
2. CPS Position Statement: Diagnosis and management of acute osteoarticular infections in children

Table 2: Transition to Oral Antibiotics
- If pathogen was obtained from culture, then oral conversion should be based on susceptibilities
- If no pathogen was obtained, then use the following for guidance:

<table>
<thead>
<tr>
<th>Empiric IV Antibiotic Used</th>
<th>Step-Down Oral Antibiotic</th>
<th>Total Duration of Antibiotic Therapy (IV and PO);</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cefazolin</td>
<td>Cephalexin</td>
<td></td>
</tr>
<tr>
<td></td>
<td>50 mg/kg/dose PO TID</td>
<td></td>
</tr>
<tr>
<td></td>
<td>≥ 40kg: 1500 mg PO qam,</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1000 mg PO mid-day, and</td>
<td></td>
</tr>
<tr>
<td></td>
<td>1500 mg PO qhs</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Dose Limit: 4g/day</td>
<td></td>
</tr>
<tr>
<td>Vancomycin</td>
<td>Consult Infectious Disease for options</td>
<td></td>
</tr>
</tbody>
</table>

4.0 Related Documents

5.0 References