**Disclaimer**

SEE ALSO: dacryocystitis, chalazion

**DESCRIPTION**

Presetable cellulitis is infection of the skin and subcutaneous tissues anterior to the orbital septum.

Orbital cellulitis (or ‘postseptal cellulitis’) is infection of the soft tissues posterior to the orbital septum, and poses risk of vision and life-threatening complications.

**BACKGROUND**

- Preseptal and orbital cellulitis occur with higher frequency in children
- Often occurs in association with sinusitis and upper respiratory tract infection (URTI)
- Pathogens: Gram positive cocci (Staphylococcus and Streptococcus species), Haemophilus species, anaerobes

**HOW TO ASSESS:**

**Red Flags:**

- Urgent surgical intervention may be required in cases of orbital cellulitis with sinusitis, subperiosteal abscess, intraorbital abscess, or foreign body. Consult oculoplastics (OPAL) and ENT.
- Intracranial infection should be suspected in patients with headache, nausea and vomiting, altered conscious state, or multiple cranial nerve palsy.
- Children with preseptal and orbital cellulitis can rapidly deteriorate. Children <4 years of age have an incomplete orbital septum and are at risk of retrograde spread of infection from the preseptal to orbital space.
- Children who are systemically unwell requiring paediatrician input may need to be transferred to The Royal Children’s Hospital (RCH). All inter-hospital transfers to and from RCH must be made consultant to consultant, involving the oculoplastics team.
- If immediate transfer/retrieval to RCH is required call ambulance or PIPER (Paediatric Infant Perinatal Emergency Retrieval - 1300 137 650)
Aetiology:

<table>
<thead>
<tr>
<th>Preseptal Cellulitis</th>
<th>Orbital Cellulitis</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Local trauma</td>
<td>• Acute sinusitis - most commonly ethmoid</td>
</tr>
<tr>
<td>• Infected chalazion</td>
<td>• Posterior extension of preseptal cellulitis</td>
</tr>
<tr>
<td>• Dacryocystitis</td>
<td>• Orbital trauma, foreign material</td>
</tr>
<tr>
<td>• URTI</td>
<td>• Dacryocystitis, dacryoadenitis</td>
</tr>
<tr>
<td>• Severe conjunctivitis</td>
<td>• Dental, facial infection</td>
</tr>
<tr>
<td>• Recent surgery on eyelids or extraocular muscles</td>
<td>• Endogenous seeding</td>
</tr>
<tr>
<td>• Endogenous seeding</td>
<td></td>
</tr>
</tbody>
</table>

On History:

- Risk factors above; ascertain immunosuppression and vaccination status (Haemophilus influenza type B (Hib))
- Acute onset of painful, swollen, red eyelids
- Orbital cellulitis: pain on eye movements, diplopia, decreased vision
- Intracranial infection: headache, nausea, vomiting, or altered conscious state

On Examination:

<table>
<thead>
<tr>
<th></th>
<th>Preseptal Cellulitis</th>
<th>Orbital Cellulitis</th>
</tr>
</thead>
<tbody>
<tr>
<td>General appearance</td>
<td>systemically well</td>
<td>may be unwell</td>
</tr>
<tr>
<td>Fever</td>
<td>variable</td>
<td>often febrile</td>
</tr>
<tr>
<td>Eyelids*</td>
<td>inflamed and swollen chalazia fluctuance - ?lid abscess</td>
<td>inflamed and swollen chalazia fluctuance - ?lid abscess</td>
</tr>
<tr>
<td>Conjunctiva</td>
<td>normal</td>
<td>chemosis</td>
</tr>
<tr>
<td>Orbital signs</td>
<td>normal</td>
<td>restricted eye movements proptosis</td>
</tr>
<tr>
<td>Optic nerve function</td>
<td>normal</td>
<td>reduced visual acuity presence of relative afferent pupil defect (RAPD) reduced colour vision reduced red saturation reduced brightness saturation abnormal visual fields</td>
</tr>
<tr>
<td>Posterior segment</td>
<td>normal</td>
<td>possible swollen optic disc choroidal folds</td>
</tr>
</tbody>
</table>

*Document dimensions of lid erythema and swelling
Differential Diagnosis:
- Severe conjunctivitis, chalazion, herpetic eye disease, dacryocystitis, dacryoadenitis, allergic dermatitis, Graves’ ophthalmopathy, idiopathic orbital inflammatory disease

On Investigation:
- Swab for microscopy, culture and viral PCR if discharge present
- **Preseptal Cellulitis:**
  - Investigations usually not necessary
- **Orbital Cellulitis suspected:**
  - Blood tests: FBE, UEC, LFT, CRP
  - Blood cultures if systemically unwell
  - CT scan (with contrast) of orbits, paranasal sinuses, and brain

Management:
- **Urgent**
  - Keep nil by mouth until assessment complete
  - Assess tetanus immunization status and administer booster if appropriate
  - Consider transfer to general hospital if patient toxic, or suspect intracranial infection:
    - Adults: discuss with St Vincent’s Hospital General Medicine Registrar, arrange ambulance transfer, consider IV antibiotics
    - Children: call ambulance or PIPER (Paediatric Infant Perinatal Emergency Retrieval - 1300 137 650) to arrange transfer/retrieval to RCH
  - Children admitted to RVEEH should be monitored using the ViCTOR track and trigger chart
- **Preseptal Cellulitis**
  - **Antibiotics**
    - **Adults**
      - Flucloxacillin 500 mg orally, 6 hourly for 7 days
      - For adults hypersensitive to penicillins (excluding immediate hypersensitivity), use:
        - Cephalexin 500 mg orally, 6 hourly for 7 days
      - For adults with immediate hypersensitivity to penicillins, use:
        - Clindamycin 450 orally, 8 hourly for 7 days
    - **Children**
      - Flucloxacillin 12.5 mg/kg (max 500 mg) orally, 6 hourly for 7 days
      - For children hypersensitive to penicillins (excluding immediate hypersensitivity), use:
        - Cephalexin 12.5 mg/kg (up to 500 mg) orally, 6 hourly for 7 days
      - For children with immediate hypersensitivity to penicillins, use:
        - Clindamycin 10 mg/kg (up to 450 mg) orally, 8 hourly for 7 days
      - If Hib infection is suspected (e.g. in unvaccinated children < 5yrs), use:
        - Amoxycillin + clavulanate 22.5+3.2 mg/kg up to 875+125 mg orally, 12 hourly for 7 days
      - For children hypersensitive to penicillins (excluding immediate hypersensitivity), use:
        - Cefuroxime:
          - 3 months – 2 years: 10 mg/kg (up to 125 mg) orally 12 hourly for 7 days
          - Older than 2 years: 15 mg/kg (up to 500 mg) orally, 12 hourly for 7 days
    - If Hib infection is suspected in patients with immediate hypersensitivity to penicillins contact the St Vincent’s Hospital Infectious Diseases team for antibiotic advice.
  - **Follow up**
    - Consider General Practitioner review in 7 days if mild disease OR
    - Acute Ophthalmology Service (AOS) in 2-7 days as clinically indicated
    - If unresponsive to oral antibiotics, or severe infection, then admit and treat as orbital cellulitis: see below
• **Orbital Cellulitis**
  
  o **Admission**
    
    Adults:
    - Admit under OPAL/ENT unit
    
    Children:
    - Admit under OPAL/ENT unit
    - Children with preseptal and orbital cellulitis can deteriorate rapidly. Children <4 years of age have an incomplete orbital septum and are at risk of retrograde extension of infection from the preseptal to orbital space.
    - Consider transfer to RCH for admission if significantly unwell, or co-morbidities (discuss with RCH team, administer IV antibiotics). Remember: all inter-hospital transfers must be made consultant to consultant involving OPAL /ENT.
  
  o **Surgical management**
    
    - OPAL and ENT teams - for consideration of surgical management of the following:
      - Extensive sinusitis with orbital cellulitis
      - Subperiosteal or intraorbital abscess
      - Intraorbital foreign body
  
  o **Antibiotics**
    
    Adults
    - Ceftriaxone 2 gram IV daily AND
    - Flucloxacillin 2 gram IV 6-hourly
    
    Children
    - Ceftriaxone 50 mg/kg (max 2g) IV daily and
    - Flucloxacillin 50 mg/kg (max 2g) IV 6-hourly
    
    - For patients with penicillin hypersensitivity contact the St Vincent’s Hospital Infectious Diseases team for antibiotic advice
    - Antibiotic selection must be modified based on results of culture and sensitivity
  
  o **Monitoring**
    
    - 4 hourly vision observations (visual acuity, pupils, colour vision)
  
  o If not improving after 24 - 48 hours of intravenous antibiotics:
    
    - Consider repeat CT to evaluate for development of abscess and need for surgical intervention
    - Discuss with St Vincent’s Hospital Infectious Diseases team for advice
  
  o If improving (e.g. afebrile with significant resolution of orbital and optic nerve signs) change to:
    
    - Amoxycillin + clavulanate 875 + 125 mg (child: 22.5 + 3.2 mg/kg up to 875 + 125 mg) orally, 12 hourly for a further 10 days.

Authors:

Dr Shivesh Varma, Dr Kristen Wells, Dr Carmel Crock and CPG Working Party

Review date:

21/08/2022
Evidence Table:

<table>
<thead>
<tr>
<th>Ref no.</th>
<th>Author</th>
<th>Title</th>
<th>Source</th>
<th>Level of evidence (I – VII)</th>
</tr>
</thead>
<tbody>
<tr>
<td>3</td>
<td>The RCH Periorbital and Orbital Cellulitis Clinical Practice Guideline</td>
<td>The RCH Periorbital and Orbital Cellulitis Clinical Practice Guideline</td>
<td>RCH Clinical Practice Guidelines 2017</td>
<td>VII</td>
</tr>
<tr>
<td>7</td>
<td>Afteh MS, Khalil HS</td>
<td>Orbital infections: five-year case series, literature review and guideline development.</td>
<td>Journal of Laryngology and Otology 2015;129:670-6</td>
<td>VI</td>
</tr>
<tr>
<td>8</td>
<td>Howe L, Jones NS</td>
<td>Guidelines for the management of periorbital cellulitis/abscess.</td>
<td>Clinical Otolaryngology &amp; Allied Sciences 2004; 29: 726-8</td>
<td>VI</td>
</tr>
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</table>

The Hierarchy of Evidence

The Hierarchy of evidence is based on summaries from the National Health and Medical Research Council (2009), the Oxford Centre for Evidence-based Medicine Levels of Evidence (2011) and Melynk and Fineout-Overholt (2011).

I Evidence obtained from a systematic review of all relevant randomised control trials.
II Evidence obtained from at least one well designed randomised control trial.
III Evidence obtained from well-designed controlled trials without randomisation.
IV Evidence obtained from well designed cohort studies, case control studies, interrupted time series with a control group, historically controlled studies, interrupted time series without a control group or with case series.
V Evidence obtained from systematic reviews of descriptive and qualitative studies.
VI Evidence obtained from single descriptive and qualitative studies.
VII Expert opinion from clinician, authorities and/or reports of expert committees or based on physiology.
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If you require clinical advice, please contact our admitting officer for assistance:
EYE: 03 9929 8033 ENT: 03 9929 8032