Pott’s puffy tumor

Brian Keehn, DO; Zachary Killeen BS; Scott A. Jorgensen, MD; Alexander J. Towbin, MD; Richard Towbin, MD.

CASE SUMMARY
A 14-year-old female presented with a 2-week history of a gradually worsening frontal headache. The patient also reported recent onset of forehead swelling.

IMAGING FINDINGS
Initial CT of the head without IV contrast demonstrated opacification of the right maxillary, right ethmoid and bilateral frontal sinuses (Figure 1). The initial CT also showed a right frontal subgaleal collection [white arrow] with an associated right frontal epidural collection [black arrow] (Figure 2). A subsequent MRI with and without contrast shows the sinus opacification [white arrows](Figure 3) with right frontal subgaleal collection and right frontal epidural abscess [*] (Figure 4).

DIAGNOSIS
Pott’s puffy tumor with intracranial extension (epidural abscess).

Differential diagnosis includes primary osseous tumor, Langerhan’s cell histiocytosis, and trauma.

DISCUSSION
English surgeon Sir Percival Pott first described Pott’s puffy tumor (PPT) in 1760. Dr. Pott had a special interest in cancer epidemiology and is also credited with describing Pott’s disease, Pott’s fracture, and various malignancies. Although first described secondary to head trauma, PPT is usually a complication of acute sinusitis, usually the frontal sinuses, characterized by a subgaleal abscess via anterior extra sinus extension. When extension of the sinus infection occurs posteriorly, an epidural abscess, subdural empyema, intracerebral abscess or dural sinus thrombosis can occur. In addition to infection eroding through the bony sinuses, direct extension secondary to trauma or surgery may be a causative source.

In the pediatric population PPT usually presents in teenagers or children with pneumatized frontal sinuses. The most common presenting symptoms of PPT include headache, fever, vomiting, forehead swelling with tenderness (hence the term “puffy”), peri orbital swelling, purulent nasal drainage and mental status changes. Although this entity is relatively uncommon, the frequency of reported cases has been increasing recently in the medical literature. As such, clinicians should be aware of these manifestations in order to accurately and appropriately treat this patient population. Age demographics are reported at a wide range from the first through eighth decades of life, though teenagers and adolescents are predominately affected and are considered the more commonly affected population for PPT.

Sinus cultures are often obtained intraoperatively and are frequently polymicrobial in nature. Organisms include alpha and beta hemolytic Streptococci, Haemophilus influenzae, Streptococcus pneumoniae, Staphylococcus, as well as less commonly Proteus, Pseudomonas, Fusobacterium and Bacteroides.
PEDIATRIC RADIOLOGICAL CASE

may be suspected following the afore-
mentioned clinical exam findings; how-
ever, neuroimaging should be con-
sidered to exclude intracranial com-
pliation. CT is the diagnostic study of choice, although MRI with contrast may help to further refine the extent of intracranial disease.

PPT is considered a surgical emer-
gency and as such should be treated accordingly. Endoscopic treatment is the preferred modality and may be performed with minimal invasiveness. Management includes abscess drainage, sequestrectomy of affected tissues and bone, as well as broad-spectrum postoperative intravenous antibiotic use for 6-8 weeks. Overall, the prognosis is generally good given an accurate diagnosis and prompt treatment.

CONCLUSION

PPT is a relatively uncommon dis-
ease that most often presents in teenag-
ers and adolescents as headache, fever, vomiting, forehead swelling with tender-
ness, periorbital swelling, purulent nasal drainage and mental status changes. These clinical exam findings, coupled with neuroimaging such as CT and MRI, may help to better define the disease process and in turn outline possible treatment approaches. Given the potential severity of PPT, prompt surgical inter-
vention and management is necessary to avoid potential long-term sequelae in these patients. Prognosis is generally good when the diagnosis is made quickly and accurately; followed by abscess drainage, bone sequestrectomy and broad-spectrum antibiotic use.

REFERENCES


Prepared by Dr. Keehn, Dr. Jorgensen and Dr. Richard Towbin while practicing at Phoenix Children’s Hospital, Phoenix, AZ; Mr. Killeen while a student at the University of Arizona College of Medicine, Phoenix, AZ; and Dr. Alexander Towbin while practicing at Cincinnati Children’s Hospital, Cincinnati, OH.