Garré’s Sclerosing Osteomyelitis with Perimandibular Soft Tissue Inflammation and Fistula

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Abstract
Garré’s sclerosing osteomyelitis is a form of chronic osteomyelitis that commonly affects children and young adults. Here, we report one such case of Garré’s sclerosing osteomyelitis in a 20-year-old female who presented with facial asymmetry and inability to open mouth. On clinical examination, it was bony hard swelling with trismus. History of infected second molar tooth extraction was present. Computed tomography scan showed thickening and sclerosis of the ramus and condylar process of mandible, on right side, with proliferative periostitis. Magnetic resonance imaging showed soft tissue edema and inflammation, in the form of enlargement of right masseter and pterygoid muscles with intramuscular fluid collection. On the basis of history, clinical signs, and imaging features, diagnosis of Garré’s osteomyelitis with fascial space infection was made. To our knowledge, very few cases of Garré’s osteomyelitis present with superimposed fascial space infection, as it is otherwise a nonsuppurative condition. Fistula formation is a very rare incidence as it is seen in our case.

Keywords ► CT proliferative periostitis ► MRI Garré’s osteomyelitis ► suppurative osteomyelitis

Introduction
Chronic osteomyelitis with proliferative periostitis (also known as periostitis ossificans or Garré’s sclerosing osteomyelitis) is a distinct form of chronic osteomyelitis.1 It is commonly associated with odontogenic infections in children and young adults.1,2 The first case was reported in tibia2 and Berger described this condition to affect mandible for the first time.1 The common causative pathogens of this condition are Staphylococci, Klebsiella, and Streptococci.2 The underlying pathology is the periosteal reaction to inflammation and hence, precisely known as chronic osteomyelitis with proliferative periostitis.1

This condition is usually nonsuppurative and asymptomatic without signs of inflammation.2-4 The SAPHO—synovitis, acne, pustulosis, hyperostosis, osteitis—syndrome is characterized by the presence of osteomyelitis in other bones, arthritis, and skin diseases.3 However, rare instances of this condition, resulting in abscess and fistula formation, are described in literature.3,4 We report one such rare case of Garré’s sclerosing osteomyelitis associated with unusual surrounding soft tissue inflammation and intramuscular abscess and fistula to skin.

Case History
A 20-year-old girl presented with swelling in the right inferior border of the mandible. On clinical examination, diffuse and tender swelling was noted, which was hard in consistency. The
The patient also had severe trismus and was unamenable for oral examination. She gave history of toothache before the onset of symptoms. Intraoral examination showed caries in right second molar tooth. Contrast-enhanced computed tomography (CECT) was advised. The imaging findings in CECT include swollen and edematous right masseter and pterygoid muscles (Fig. 1). There was intramuscular fluid collection within the masseter muscle and fistulous communication to skin (Fig. 2). Three-dimensional computed tomography (CT) of mandible showed cortical thinning involving ramus and condylar process of mandible, on right side, characteristic of Garré’s sclerosing osteomyelitis (Fig. 3). Magnetic resonance imaging (MRI) also showed hyper-intensities in right masseter and pterygoid muscles, which implicates inflammation (Fig. 4). Incision and drainage of the collection, followed by biopsy from expanded cortical regions, was done and the patient was put on intravenous antibiotics.

Discussion

Garré’s sclerosing osteomyelitis was first described by Carl Garré in 1893 in tibia, which resulted from radiation exposure. However, it was Berger who first described this condition affecting the jaw bones. This chronic form of osteomyelitis is usually asymptomatic without any signs of local inflammation. The common organisms encountered in the disease process include Staphylococcus, Klebsiella, and Streptococcus, resulting in phases of remission and exacerbation. The commonest cause is odontogenic infection but it can also occur in gunshot wounds, fractures, pyoderma, postoperative bone interventions, etc. The severity and duration of disease depends on many factors like the virulence of the causative organisms, the presence of underlying diseases, and the immunity of the host. It is often unilateral and nonsuppurative. However, rare instances of this condition, resulting in abscess and fistula formation, are described in literature.

Facial asymmetry is often the presenting complaint and pain is not a characteristic finding. The other markers of acute inflammation like fever, white blood cell count, and C-reactive protein may also not elevate characteristically. The disease process starts in the spongiosa and extends into periosteum, resulting in osteoblastic reaction. Unfortunately in some patients the disease process can further extend to the perimandibular soft tissue with resultant abscess and fistula formation. In such a scenario, there can be severe trismus resulting from the masticator space infection.
Imaging plays a very important role in the diagnosis of this condition. The CT imaging features of periostitis ossificans include cortical thinning and periosteal thickening with lamellar appearance (onion skin), commonly affecting the ramus of mandible. The laminated appearance is due to modulation of fibroblasts in the adjacent soft tissue, which develop osteoblastic capacity and give rise to sheets of new bone. The differential diagnosis of this type of periostitis includes Caffey disease, Ewing’s sarcoma, osteosarcoma, fibrous dysplasia, osteoma, exostosis, and ossifying subperiosteal hematoma. Caffey disease or infantile cortical hyperostosis is a rare self-limiting condition of infancy that is characterized by cortical hyperostosis, particularly affecting the mandible and facial bones. This condition is bilateral and multiple bones are involved unlike Garré’s osteomyelitis. Ewing’s sarcoma and osteosarcoma are the two malignant conditions with similar periosteal reaction, although they are very rare in mandible and characterized by “sun ray” appearance. While the latter is characterized by Codman triangle, the former, in addition, shows osteolytic areas and neurological symptoms like facial neuralgia and lip paresthesia.

Fibrous dysplasia is typically characterized by the “ground glass appearance” and the enlargement is seen in the bone matrix, whereas in Garre’s osteomyelitis it is seen on the outer surface of the cortex. The appearance of ossifying subperiosteal hematoma or fracture with callus formation may mimic Garre’s osteomyelitis clinically. However, the former does not exhibit uniform radiopacity and display mottled appearance or trabecular structure while the absence of trauma history can exclude both the conditions.

Once the diagnosis of Garre’s osteomyelitis is made, the most commonly accepted treatment is the administration of antibiotics and the extraction of the infected tooth. However, when complicated with abscess formation as demonstrated in CECT as peripherally enhancing fluid collection, it requires incision and drainage. MRI is helpful in presupplicative stages, when muscle edema is detected as increased signals within the muscles in the T2 and short tau inversion recovery sequences. It is important to identify the perimandibular soft tissue infection, as otherwise conservative therapy and removing the causative factor are usually sufficient for Garre’s osteomyelitis.

Declaration of Patient Consent
The authors certify that they have obtained all appropriate patient consent forms. In the form the patient(s) has/have given his/her/their consent for his/her/their images and other clinical information to be reported in the journal. The patients understand that their names and initials will not be published and due efforts will be made to conceal their identity, but anonymity cannot be guaranteed.

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Conflicts of Interest
There are no conflicts of interest.

References